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(21) International Application Number: PCT/US97/19851 (22) International Filing Date: 31 October 1997 (31.10.97) (30) Priority Data: 60/030,367 5 November 1996 (05.11.96) US (71) Applicant: BRISTOL-MYERS SQUIBB COMPANY [US/US]; P.O. Box 4000, Princeton, NJ 08543-4000 (US). (72) Inventors: KING, Dalton; 114 Wakefield Street, Hamden, CT 06514 (US). FIRESTONE, Raymond, A.; 387 Temple Street, New Haven, CT 06511 (US). DUBOWCHIK, Gene, M.; 65 Spring Street, Middlefield, CT 06455 (US). (74) Agents: SAVITSKY, Thomas, R. et al.; Bristol-Myers Squibb Company, P.O. Box 4000, Princeton, NJ 08543-4000 (US).		(81) Designated States: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: BRANCHED PEPTIDE LINKERS (57) Abstract Conjugates containing a targeting ligand, such as an antibody, a therapeutically active drug and a branched peptide linker. The branched peptide linker contains two or more amino acid moieties that provide an enzyme cleavage site. The number of drugs capable of being bonded to the branched linkers varies by a factor of two for each generation of branching.		

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BRANCHED PEPTIDE LINKERS**Background of the Invention**

Bifunctional compounds which link cytotoxic reagents to antibodies
5 (i.e., "linkers") are known in the art. These compounds have been particularly useful in the formation of immunoconjugates directed against tumor associated antigens. Such immunoconjugates allow the selective delivery of toxic drugs to tumor cells. (See e.g., Hermentin and Seiler, "Investigations With Monoclonal Antibody Drug Conjugates," Behring Insti.
10 Mitl. 82:197-215 (1988); Gallego et al., "Preparation of Four Daunomycin-Monoclonal Antibody 791T/36 Conjugates With Anti-Tumor Activity". Int. J. Cancer 33:737-44 (1984); Arnon et al., "In Vitro and In Vivo Efficacy of Conjugates of Daunomycin With Anti-Tumor Antibodies," Immunological Rev. 62:5-27 (1982).

15 Greenfield et al. have described the formation of acid-sensitive immunoconjugates containing the acylhydrazide conjugated via an acylhydrazone bond to the 13-keto position of an anthracycline molecule, and conjugation of this anthracycline derivative to an antibody molecule (Greenfield et al., European Patent Publication EP 0 328 147, published
20 August 16, 1989, which corresponds to pending U.S. Serial No. 07/270,509, filed November 16, 1988 and U.S. Serial No. 07/155,181, filed February 11, 1988, now abandoned). This latter reference also discloses specific thioether-containing linkers and conjugates, including hydrazone thioether containig immunoconjugates.

25 Kaneko et al. (U.S. Serial No. 07/522,996, filed May 14, 1990, which is equivalent to European Patent Publication, EP A 0 457 250, published November 21, 1991) have also described the formation of conjugates containing anthracycline antibiotics attached to a bifunctional linker by an acylhydrazone bond at the C-13 position of an anthracycline
30 molecule. In their invention the linkers contain a reactive pyridinyldithio- or an ortho-nitrophenyldithio- group, by which the linker reacts with a suitable group attached to a cell reactive ligand, to form the completed conjugate.

Conjugates which rely on simple acid hydrolysis may release the drug prematurely. Accordingly, it would be desirable to have conjugates that release active drug in a more site-specific fashion. European Patent Publication 94107501.2 discloses lysosomal enzymes-cleavable antitumor drug conjugates which are selectively activatable at the site of the tumor. However, one of the problems with prior art immunoconjugates is the relatively low ratio of drug to targeting ligand (e.g., immunoglobulin) achievable. It would be highly desirable to have immunoconjugates, activatable at the tumor site, which provide a higher ratio of drug to targeting ligand.

Summary of the Invention

The present invention provides novel branched peptide linkers. The novel linkers are used to prepare novel drug/linker molecules and biologically active conjugates composed of a targeting ligand, a therapeutically active drug, and a branched peptide linker. The novel conjugates are selectively activatable at the site of of a selected target cell population recognized by the targeting ligand.

As used herein the term "drug/linker" or "linker/drug" refers to the branched peptide linker molecule coupled to two or more therapeutically active drug molecules, and the term "conjugate" refers to the drug/linker molecule coupled to the targeting ligand.

The branched peptide linker contains a protein peptide spacer and may also contain a self-immolating spacer which spaces the protein peptide sequence and the drug. The linkers of the invention are branched so that more than one drug molecule per linker are coupled to the targeting ligand. The number of drugs attached to each linker varies by a factor of 2 for each generation of branching. Thus, the number of drug molecules per molecule of linker can be 2, 4, 8, 16, 32, 64, etc. The factor of branching can be expressed mathematically as 2^n , wherein n is a positive integer. Thus, a singly branched linker will have a first generation of branching or 2^1 , i.e., contains a potential of two drug molecules per linker. A doubly branched

linker will have a second generation of branching or 2^2 , i.e., contains a potential of four drug molecules per linker.

As n rises from $n=1$, there is a tendency for the solubility of the immunoconjugate to diminish. Solubility can be enhanced by using more water-soluble peptides, or addition of a water-solubilizing moiety such as polyethylene glycol or charged species, e.g., β -alanine, to the drug in such a way that it is released from the drug by either the low pH or the enzymes of the liposomal milieu

The present invention is directed to a branched peptide linker for linking a thiol group derived from a targeting ligand to two or more drug moieties which comprises a compound having a terminus containing a thiol acceptor for binding to a thiol group (also called a sulfhydryl group) derived from a targeting ligand, at least one point of branching which is a polyvalent atom, such as a carbon atom or a nitrogen atom, allowing for a level of branching of 2^n , wherein n is a positive integer, at least two amino acid moieties per branch providing at least one enzymatic site per branch, and at least two other termini containing groups capable of forming covalent bonds with chemically reactive functional groups derived from a drug moiety. It is preferred that n is 1, 2, 3, or 4; more preferably 1, 2, or 3; and most preferably 1 or 2. It is also preferred that the targeting ligand is an antibody or fragment thereof.

As used in the preceeding paragraph, the phrase "thiol group derived from the targeting ligand" means that the thiol group is already present on the targeting ligand or that the targeting ligand is chemically modified to contain a thiol group, which modification optionally includes a thiol spacer group between the targeting ligand and the thiol group. Likewise, the phrase "chemically reactive functional group derived from a drug moiety" means that the chemically reactive functional group is already present on the drug or the drug is chemically modified to contain such chemically reactive functional group. Such chemically reactive functional groups are groups that are capable of forming covalent bonds with a linker terminus. Examples of such chemically reactive functional groups include

primary or secondary amino, hydroxyl, sulfhydroxyl, carboxyl, aldehyde, ketone, and the like.

Also provided by the invention are intermediates for preparing the linkers, drug/linkers, and/or conjugates; and a method for treating or preventing a selected disease state which comprises administering to a patient a conjugate of the invention.

The selected target cell population recognized by the targeting ligand is preferably a tumor.

An aspect of the invention provides tumor-specific conjugates which are highly selective substrates for drug-activating enzymatic cleavage by one or more tumor-associated enzymes.

A further aspect of the invention provides tumor-specific drug conjugates wherein the activating enzyme is one which is present in the tumor in sufficient amounts to generate cytotoxic levels of free drug in the vicinity of the tumor.

Another aspect of the invention provides tumor-specific drug conjugates which are stable to adventitious proteases in blood.

A still further aspect of the present invention provides a tumor-specific conjugate in accordance with the preceding aspects, which is considerably less toxic than the activated drug.

In another aspect the present invention provides methods for delivering the conjugates to target cells in which a modification in biological process is desired, such as in the treatment of diseases such as cancer.

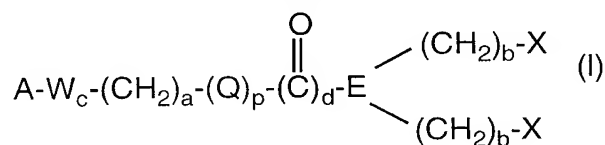
The present invention also provides a method for delivering to the site of tumor cells in a warm-blooded animal an active antitumor drug by administering to said warm-blooded animal the conjugate according to this invention.

The above and other aspects of the present invention are achieved by derivatizing a drug (e.g., an antitumor agent) linked to a ligand through a peptide linker, made up of a protein peptide sequence and a self-immolating spacer, at a reactive site appropriate for inhibiting the pharmacological activity of the antitumor agent to thereby convert the antitumor agent into a pharmacologically inactive peptidyl derivative conjugate. The peptide linker has at least two amino acid residue

sequences specifically tailored so as to render the peptidyl derivatives selective substrates for drug-activating enzymatic cleavage by one or more lysosomal proteases, such as cathepsin B, C, D, or L. The enzymatic cleavage reaction will remove the peptide linker moiety from the drug conjugate and effect release of the antitumor agents in pharmacologically active form selectively at the tumor site. In comparison with ligand-drug linkers which rely on simple acid hydrolysis for drug release this new method provides significantly less systemic toxicity due to premature linker hydrolysis in the blood, consequently a greater amount of the drug is delivered to the tumor site, and the method results in a longer storage life and simplified handling conditions for the conjugate.

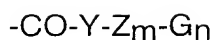
The conjugates of the present invention show significantly less systemic toxicity than biparte conjugates and free drug. The conjugates of the invention retain both specificity and therapeutic drug activity for the treatment of a selected target cell population. They may be used in a pharmaceutical composition, such as one comprising a pharmaceutically effective amount of a compound of Formula (II) below, associated with a pharmaceutically acceptable carrier, diluent or excipient.

The present invention is directed to a linker molecule of the formula



wherein

- A is a thiol acceptor;
- W is a bridging moiety;
- c is an integer of 0 to 1;
- a is an integer of 2 to 12;
- Q is O, NH, or N-lower alkyl;
- p is an integer of 0 or 1;
- d is an integer of 0 or 1;
- E is a polyvalent atom;
- each b is an integer of 1 to 10;
- each X is of the formula



wherein

Y is two amino acid residues in the L form;

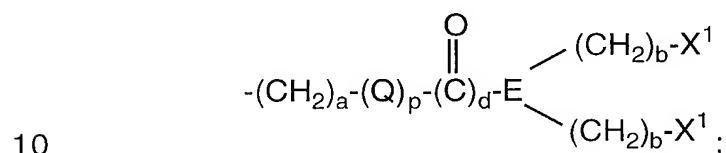
Z is one or two amino acid residues;

5 m is an integer of 0 or 1;

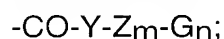
G is a self-immolative spacer; and

n is a integer of 0 or 1; provided that when G is 0 then $-\text{Y}-\text{Z}_m$ is ala-leu-ala-leu or gly-phe-leu-gly;

or each X is of the formula

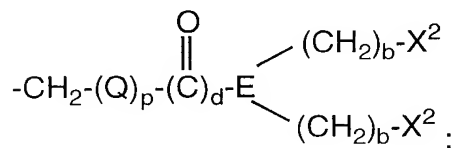


wherein each X^1 is of the formula

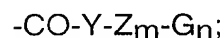


and wherein Y, Z, Q, E, G, m, d, p, a, b and n are as defined above;

15 or each X^1 is of the formula



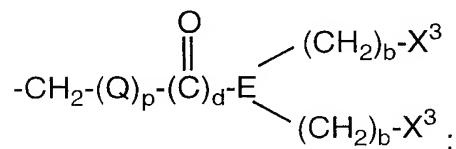
wherein each X^2 is of the formula



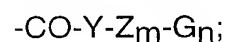
and wherein Y, Z, G, Q, E, m, d, p, a, b and n are as defined above;

20

or each X^2 is of the formula

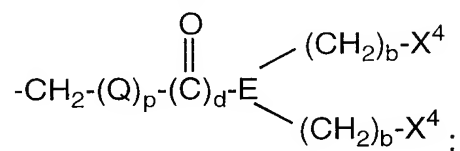


wherein each X^3 is of the formula



and wherein Y, Z, G, Q, E, m, d, p, a, b and n are as defined above;

or each X³ is of the formula

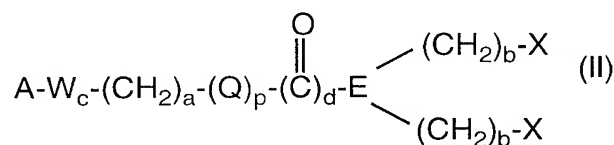


5 wherein each X⁴ is of the formula

-CO-Y-Z_m-G_n;

and wherein Y, Z, G, Q, E, m, d, p, a, b and n are as defined above.

10 The present invention is also directed to a drug/linker molecule of the formula



wherein

A is a thiol acceptor;

W is a bridging moiety;

15 c is an integer of 0 to 1;

a is an integer of 2 to 12;

Q is O, NH, or N-lower alkyl;

p is an integer of 0 or 1;

d is an integer of 0 or 1;

20 E is a polyvalent atom;

each b is an integer of 1 to 10;

each X is of the formula

-CO-Y-Z_m-G_nD

wherein

25 Y is two amino acid residues in the L form;

Z is one or two amino acid residues;

m is an integer of 0 or 1;

G is a self-immolative spacer;

n is a integer of 0 or 1; provided that when G is 0 then -Y-Z_m-

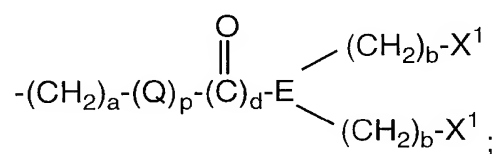
is ala-leu-ala-leu or gly-phe-leu-gly; and

D is a Drug moiety having a backbone and at least one chemically reactive functional group pendant thereto

5 chemically

reacted to the self-immolative spacer or terminal amino acid residue to form a covalent bond, said functional group selected from the group consisting of a primary or secondary amine, hydroxyl, sulfhydryl, carboxyl, aldehyde or ketone;

10 or each X is of the formula

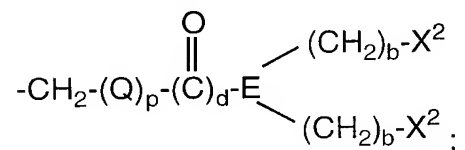


wherein each X¹ is of the formula

-CO-Y-Z_m-G_n-D;

wherein Y, Z, G, D, Q, E, m, d, p and n are as defined above;

15 or each X¹ is of the formula



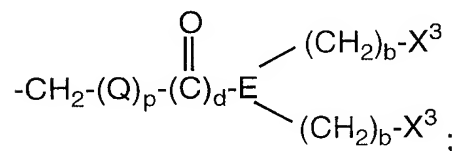
wherein each X² is of the formula

-CO-Y-Z_m-G_n-D;

and wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined above;

20

or each X² is of the formula

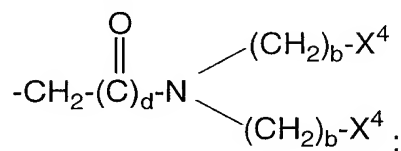


wherein each X³ is of the formula

-CO-Y-Z_m-G_n-D;

and wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined above;

or each X³ is of the formula



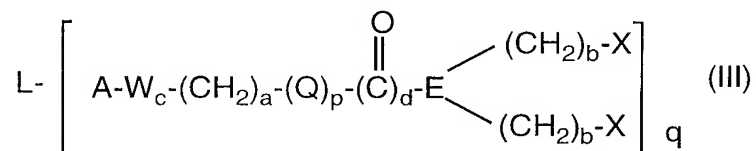
5 and wherein each X⁴ is of the formula

-CO-Y-Z_m-G_n-D;

and wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined above.

Furthermore, the present invention is directed to conjugate of the

10 formula



wherein

L is a ligand;

q is an integer of 1 to 10;

15 A is a thiol acceptor;

W is a bridging moiety;

c is an integer from 0 to 1;

a is an integer of 2 to 12;

Q is O, NH, or N-lower alkyl;

20 p is an integer of 0 or 1;

d is an integer of 1 or 2;

E is a polyvalent atom;

each b is an integer of 1 to 10;

each X is of the formula

25 -CO-Y-Z_m-G_n-D

wherein

Y is two amino acid residues in the L form;

Z is one or two amino acid residues;

m is an integer of 0 or 1;

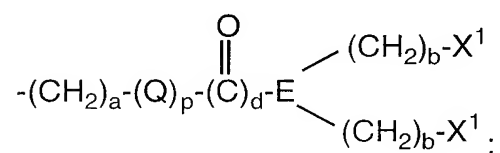
G is a self-immolative spacer; and

n is a integer of 0 or 1; provided that when G is 0 then -Y-Z_m- is ala-leu-ala-leu or gly-phe-leu-gly;

5 D is a drug moiety having a backbone and at least one chemically reactive functional group pendant thereto reacted to the self-immolative spacer to form a covalent bond, said functional group selected from the group consisting of a primary or secondary amine, hydroxyl, carboxyl, sulfhydryl, aldehyde, ketone;

10 or

or X is of the formula

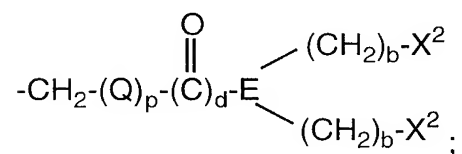


wherein each X¹ is of the formula

-CO-Y-Z_m-G_n-D;

15 and wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined above;

or each X¹ is of the formula

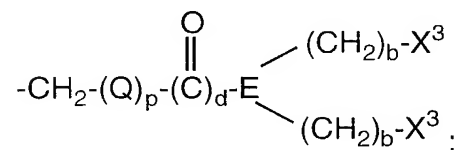


wherein each X² is of the formula

20 -CO-Y-Z_m-G_n-D;

and wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined above;

or each X² is of the formula

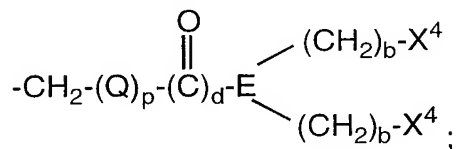


25 wherein each X³ is of the formula

-CO-Y-Z_m-G_n-D;

wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined above;

or each X³ is of the formula



5

wherein end X⁴ is of the formula

-CO-Y_m-Z_m-G_n-D;

wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined above.

10 As used herein the term "lower alkyl" is an alkyl group having 1 to 3 carbon atoms. It is preferred that the polyvalent atom is carbon or nitrogen.

In one embodiment the drug moiety is an anthracycline antibiotic and the targeting ligand is an antibody.

15 In a preferred embodiment the anthracycline is bound to the linker at an amino sugar group of the anthracycline. It is preferred that the sugar moiety is daunosamine. The antibody then is bound, through the linker, to the anthracycline compound. In an especially preferred embodiment, this linkage occurs through a reduced disulfide group (i.e. a free sulfhydryl group (-SH)) on an antibody).

20 In a most preferred embodiment the anthracycline drug moiety is doxorubicin; the thiol acceptor is a Michael Addition acceptor (from which a Michael Addition Adduct is derived), more preferably is a maleimido- group; and the antibody moiety is a chimeric or humanized antibody, and the point of attachment of the linker to the drug is at the amino group of the sugar

25 moiety of the drug.

The conjugates of the invention retain both specificity and therapeutic drug activity for the treatment of a selected target cell population. They may be used in a pharmaceutical composition, such as one comprising a pharmaceutically effective amount of a compound of Formula (III) associated

30 with a pharmaceutically acceptable carrier, diluent or excipient.

Detailed Description of the Invention

The present invention provides novel drug-linker-ligand conjugates composed of a ligand capable of targeting a selected cell population, and a drug connected to the ligand by a branched peptide linker. The linker contains a thiol acceptor such as a Michael Addition acceptor, a bridging moiety, a point of branching, and a peptide sequence per each branch containing at least two amino acid moieties which provides an enzymatic cleavage site for an enzyme such as cathepsin B, C, D, or L. The branched peptide linker may also contain a self-immolating spacer, which spaces the drug and the protein peptide sequence. It has been discovered that when the peptide sequence (i.e., Y-Z_m) is ala-leu-ala-leu- or gly-phe-leu-gly, then the presence of the self-immolative spacer is optional. The reason that a self-immolative spacer is not required for these peptide sequences is not entirely understood; however, it may be due to response to different enzymes.

The targeting ligand molecule can be an immunoreactive protein such as an antibody, or fragment thereof, a non-immunoreactive protein, or peptide ligand such as bombesin or, a binding ligand recognizing a cell associated receptor such as a lectin, or any protein or peptide that possesses a reactive sulfhydryl group (-SH) or can be modified to contain such a sulfhydryl group. The thiol acceptor carboxylic is linked to the ligand via a thioether bond, and the drug is linked to the linker via a functional group selected from primary or secondary amine, hydroxyl, sulfhydryl, carboxyl, aldehyde or ketone.

For a better understanding of the invention, the drugs, ligands, peptides and spacers will be discussed individually. The synthesis of the conjugates then will be explained.

It will be understood that in the following detailed description and appended claims, the abbreviations and nomenclature employed are those which are standard in amino acid and peptide chemistry, and that all the amino acids referred to are in the L-form unless otherwise specified.

Some abbreviations used in the present application, unless otherwise indicated, are as follows:

AcOH: acetic acid; Alloc: allyloxy-carbonyl; Boc: t-butyloxycarbonyl; DBU: 1,4-diazobicycloundecene; DCC: dicyclohexylcarbodiimide; DCI: direct chemical ionization; DCU: dicyclohexylurea; DIEA: diisopropylethylamine; DMAP: 4-dimethylaminopyridine; DME: 1,2-dimethoxyethane; DOX: doxorubicin; DTT: dithiothreitol; EEDQ: N-ethoxycarbonyl-2-ethoxy-1,2-dihydroquinoline; EtOAc: ethyl acetate; FAB: fast atom bombardment; Fmoc: fluorenylmethoxycarbonyl; GABA: γ -aminobutyric acid; HOBt: N-hydroxybenzotriazole; HRMS: high resolution mass spectroscopy; LDL: low density lipoprotein; MC: 6-maleimidocaproyl; MP: 3-maleimidopropionyl; MPr-BHP: maleimidopropyl-bis-hydroxypropyl; MPr-Mal: maleimidopropyl-malonyl; MEt-IBHE: maleimidoethyl-imino-bis-hydroxyethyl; MMA: mitomycin A, MMC: mitomycin C; Mtr: 4-methoxytrityl; NHS: N-hydroxysuccinimide; NMP: N-methylpyrrolidinone; PABC: p-aminobenzyl-carbamoyl; PAB-OH: p-aminobenzyl alcohol; PNP: p-nitrophenol; TFA: trifluoroacetic acid; THF: tetrahydrofuran.

The Peptide Residue (-Y-Z_m-)

The peptide linker of the present invention contains two, three or four amino acid residues per branch, together with the self-immolative spacer if present, that provides one or more enzyme cleavage sites. The amino acid moieties collectively form a peptide sequence.

The amino acid residues making up the peptide residues are each selected, independently, from the group of amino acids, preferably naturally occurring amino acids. The naturally occurring amino acids are alanine (Ala), arginine (Arg), asparagine (Asn), aspartic acid (Asp), cysteine (Cys), glutamic acid (Glu), glutamine (Gln), glycine (Gly), histidine (His), isoleucine (Ile), leucine (Leu), lysine (Lys) methionine (Met), ornithine (Orn), phenylalanine (Phe), proline (Pro), serine (Ser), threonine (Thr), tryptophan (Trp), tyrosine (Tyr), and valine (Val). Preferred naturally occurring amino acids are Ala, Val, Leu, Lys, Ile, Met, Phe, Trp, and Pro. Certain non-naturally occurring amino acids also can be part of the peptide residue. Such non-naturally occurring amino acids include citrulline (Cit) and protected amino acids such as naturally occurring amino acids protected with groups such as acetyl, formyl, tosyl, nitro and the like. When specific amino acids

are indicated herein as part of a peptide sequence, they are in the L form unless specified otherwise. The amino acid residues making up the "Y" moiety must be in the L form. The amino acid residue(s) making up the "Z" moiety can be either in the L or D form. More preferred amino acids include Lys, Lys protected with acetyl or formyl, Arg, Arg arginine protected with tosyl or nitro groups, His, Orn, Orn protected with acetyl or formyl, Phe, Val, Ala, and Cit. Most preferred are Lys and Cit..

The amino acid residue sequence is specifically tailored so that it will be selectively enzymatically cleaved from the resulting peptidyl derivative drug-conjugate by one or more of the tumor-associated proteases.

The amino acid residue chain length of each branch of the peptide linker preferably ranges from that of a dipeptide to that of a tetrapeptide.

It is preferred that the first amino acid residue (i.e., the first amino acid making up "Y", read left to right) is a basic amino acid (e.g., Lys or Arg) or has strong hydrogen bonding capability (e.g., Cit). It is preferred that the second amino acid residue (i.e., the second amino acid making up "Y", read left to right) has a hydrophobic side chain (e.g., Phe, Val, Ala, Leu or Ile).

The following group of exemplary peptide linker groups, are named in order to illustrate further the conjugates of the present invention:

Phe-Lys, Val-Lys, Phe-Phe-Lys, Lys-Phe-Lys, Gly-Phe-Lys, Ala-Lys, Val-Cit, Phe-Cit, Leu-Cit, Ile-Cit, Trp-Cit, Phe-Ala, Gly-Phe-Leu-Gly, Ala-Leu-Ala-Leu, Phe-Ng-tosyl-Arg, Phe-Ng-Nitro-Arg, Lys-Lys, Lys-Cit, and Cit-Cit.

Specific examples of the preferred embodiment of peptide sequences include Phe-Lys, Val-Lys, Val-Cit, and D-Phe-Phe-Lys.

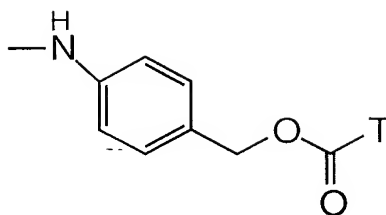
Numerous specific peptide linker molecules suitable for use in the present invention can be designed and optimized in their selectivity for enzymatic cleavage by a particular tumor-associated protease. The preferred peptide linkers for use in the present invention are those which are optimized toward the protease, cathepsin B, C, D, and L.

The Spacer ("G")

The molecules in accordance with the present invention may employ an intermediate self-immolative spacer moiety which spaces and covalently links together the drug moiety and the protein peptide moiety. A self-immolative spacer may be defined as a bifunctional chemical moiety which is capable of covalently linking together two spaced chemical moieties into a normally stable tripartate molecule, releasing one of said spaced chemical moieties from the tripartate molecule by means of enzymatic cleavage; and following said enzymatic cleavage, spontaneously cleaving from the remainder of the molecule to release the other of said spaced chemical moieties. In accordance with the present invention, the self-immolative spacer is covalently linked at one of its ends to the protein peptide moiety and covalently linked at its other end to the chemical reactive site of the drug moiety whose derivatization inhibits pharmacological activity, so as to space and covalently link together the protein peptide moiety and the drug moiety into a tripartate molecule which is stable and pharmacologically inactive in the absence of the target enzyme, but which is enzymatically cleavable by such target enzyme at the bond covalently linking the spacer moiety and the protein peptide moiety to thereby effect release of the protein peptide moiety from the tripartate molecule. Such enzymatic cleavage, in turn, will activate the self-immolating character of the spacer moiety and initiate spontaneous cleavage of the bond covalently linking the spacer moiety to the drug moiety, to thereby effect release of the drug in pharmacologically active form.

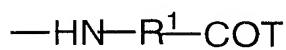
In the molecules of Formulas I, II, and III:

G is a self-immolative spacer moiety which spaces and covalently links together the drug moiety and the amino acid, in which the spacer is linked to the drug moiety via the T moiety (as used in the following formulas "T" represents a nucleophilic atom which is already contained in the Drug), and which may be represented by the structures of Formulae, (IV), (V), (VI), (VII), or (VIII):



Formula (IV)

in which T is O, N or S,

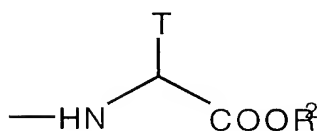


Formula (V)

5

in which T is O, N or S, and

R¹ is C₁-C₅ alkyl;

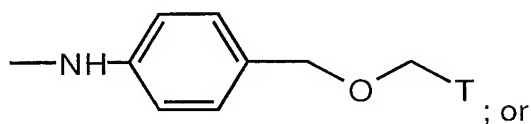


Formula (VI)

10 (J. Med. Chem., 27: 1447 (1984)

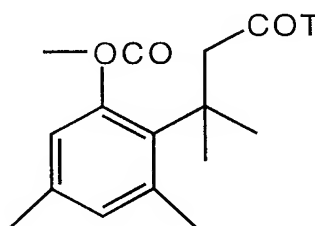
in which T is O, N or S, and

R² is H or C₁-C₅ alkyl;



Formula (VII)

15 in which T is O, N or S,



Formula (VIII)

in which T is O, S or N.

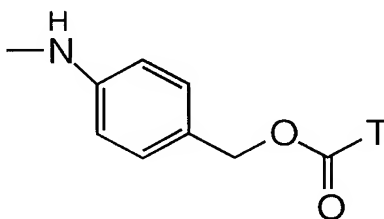
20

As used herein "C₁-C₅ alkyl" is meant to include a branched or unbranched hydrocarbon chain having, unless otherwise noted, one to five

carbon atoms, including but not limited to methyl, ethyl, isopropyl, n-propyl, sec-butyl, isobutyl, n-butyl and the like.

A preferred G self-immolative spacer moiety suitable for use in the present invention is PABC represented by the Formula (IVa):

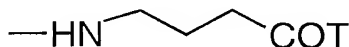
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Formula (IVa)

Another preferred G self-immolative spacer moiety suitable for use in the present invention is GABA represented by the Formula (Va):

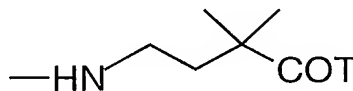
10



Formula (Va)

Yet another preferred G self-immolative spacer moiety suitable for use in the present invention is α,α -dimethyl GABA represented by the Formula (Vb):

15



Formula (Vb)

Another preferred G self-immolative spacer moiety suitable for use in the present invention is β,β -dimethyl GABA represented by the Formula (Vc):

20



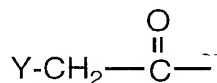
Formula (Vc)

The Thiol Acceptor

25

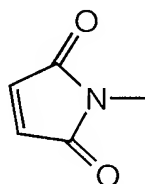
In the molecules of Formulas I, II, and III, the thiol acceptor "A" is linked to the ligand via a sulfur atom derived from the ligand. The thiol acceptor becomes a thiol adduct after bonding to the ligand through a thiol group via

a thioester bond. The thiol acceptor can be, for example, an alpha-substituted acetyl group. Such a group has the formula

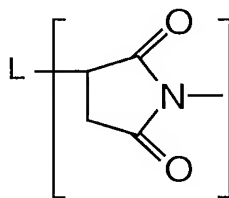


wherein Y is a leaving group. Examples of leaving groups include Cl, Br, I, mesylate, tosylate, and the like. If the thiol acceptor is an alpha-substituted acetyl group, the thiol adduct after linkage to the ligand forms the bond -S-CH₂-

Preferably, the thiol acceptor is a Michael Addition acceptor. A representative Michael Addition acceptor of this invention has the formula



After linkage the thiol group of the ligand, the Michael Addition acceptor becomes a Michael Addition adduct, such as of the formula A

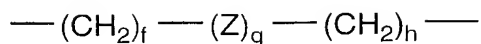


A

wherein L is ligand.

The Bridging Group ("W")

The bridging group is a bifunctional chemical moiety which is capable of covalently linking together two spaced chemical moieties into a stable tripartate molecule. Examples of bridging groups are described in S.S. Wong, Chemistry of Protein Conjugation and Crosslinking, CRC Press, Florida, (1991); and G.E. Means and R.E. Feeney, Bioconjugate Chemistry, vol. 1, pp.2-12, (1990), the disclosures of which are incorporated herein by reference. Specifically, the bridging group "W" covalently links the thiol acceptor to a keto moiety. An example of a bridging group has the formula



wherein

f is an integer of 0 to 10,

h is an integer of 0 to 10,

5 g is an integer of 0 or 1,

provided that when g is 0, then f + h is 1 to 10,

Z is S, O, NH, SO₂, phenyl, naphthyl, a polyethylene glycol, a cycloaliphatic hydrocarbon ring containing 3

to 10 carbon atoms, or a heteroaromatic

10 hydrocarbon ring containing 3 to 6 carbon

atoms and 1 or 2 heteroatoms selected from O,

N, or S.

Preferred cycloaliphatic moieties include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, and the like. Preferred heteroaromatic moieties

15 include pyridyl, polyethylene glycol (1-20 repeating units), furanyl, pyranal, pyrimidinyl, pyrazinyl, pyridazinyl, oxazinyl, pyrrolyl, thiazolyl, morpholinyl, and the like.

In the bridging group it is preferred that when g is 0, f + h is an integer of 2 to 6 preferably 2 to 4 and more preferably 2. When g is 1, it is preferred
20 that f is 0, 1 or 2, and that h is 0, 1 or 2.

Preferred bridging groups coupled to thiol acceptors are shown in the Pierce Catalog, pp. E-12, E-13, E-14, E-15, E-16, and E-17 (1992).

The Drug

25 The drug conjugates of the present invention are effective for the usual purposes for which the corresponding drugs are effective, and have superior efficacy because of the ability, inherent in the ligand, to transport the drug to the desired cell where it is of particular benefit. Further, because the conjugates of the invention can be used for modifying a given biological
30 response, the drug moiety is not to be construed as limited to classical chemical therapeutic agents. For example, the drug moiety may be a protein or polypeptide possessing a desired biological activity. Such proteins may include, for example, a protein such as tumor necrosis factor.

The preferred drugs for use in the present invention are cytotoxic drugs, particularly those which are used for cancer therapy. Such drugs include, in general, DNA damaging agents, anti-metabolites, natural products and their analogs. Preferred classes of cytotoxic agents include, for example, the enzyme inhibitors such as dihydrofolate reductase inhibitors, and thymidylate synthase inhibitors, DNA intercalators, DNA cleavers, topoisomerase inhibitors, the anthracycline family of drugs, the vinca drugs, the mitomycins, the bleomycins, the cytotoxic nucleosides, the pteridine family of drugs, diynenes, the podophyllotoxins, differentiation inducers, and taxanes. Particularly useful members of those classes include, for example, methotrexate, methopterin, dichloromethotrexate, 5-fluorouracil, 6-mercaptapurine, cytosine arabinoside, melphalan, leurosine, leurosideine, actinomycin, daunorubicin, doxorubicin, mitomycin C, mitomycin A, carminomycin, aminopterin, tallysomycin, podophyllotoxin and podophyllotoxin derivatives such as etoposide or etoposide phosphate, vinblastine, vincristine, vindesine, taxol, taxotere, retinoic acid, butyric acid, N⁸-acetyl spermidine, camptothecin, and their analogues.

As noted previously, one skilled in the art may make chemical modifications to the desired compound in order to make reactions of that compound more convenient for purposes of preparing conjugates of the invention.

In the conjugate of Formula I,
D is a drug moiety having pendant to the backbone thereof a chemically reactive functional group by means of which the drug backbone is bonded to the protein peptide linker, said functional group selected from the group consisting of a primary or secondary amine, hydroxyl, sulfhydryl, carboxyl, aldehyde or a ketone. Representative of said amino containing drugs are mitomycin-C, mitomycin-A, daunorubicin, doxorubicin, aminopterin, actinomycin, bleomycin, 9-amino camptothecin, N⁸-acetyl spermidine, 1-(2-chloroethyl)-1,2-dimethanesulfonyl hydraxide, tallysomycin, cytarabine and derivatives thereof.

Representative of said alcohol group containing drugs are etoposide, camptothecin, taxol, esperamicin, 1,8-dihydroxy-bicyclo[7.3.1]trideca-4-9-diene-2,6-diyne-13-one, (U.S. Patent 5,198,560), podophyllotoxin, anguidine, vincristine, vinblastine, morpholine-doxorubicin, n-(5,5-diacetoxy-pentyl)doxorubicin, and derivatives thereof.

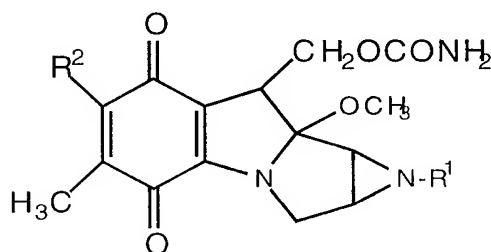
Representative of said sulfhydryl containing drugs are esperamicin and 6-mercaptopurine, and derivatives thereof.

Representative of said carboxyl containing drugs are methotrexate, camptothecin (ring-opened form of the lactone), butyric acid, retinoic acid, and derivatives thereof.

Representative of said aldehyde and ketone containing drugs are anguidine and anthracyclines such as doxorubicin, and derivatives thereof.

A highly preferred group of cytotoxic agents for use as drugs in the present invention include drugs of the following formulae:

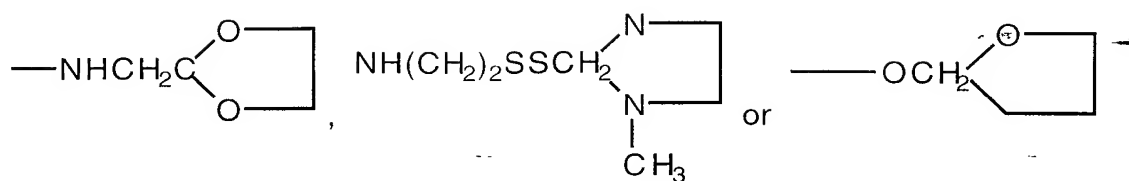
THE MITOMYCIN GROUP OF FORMULA (1):



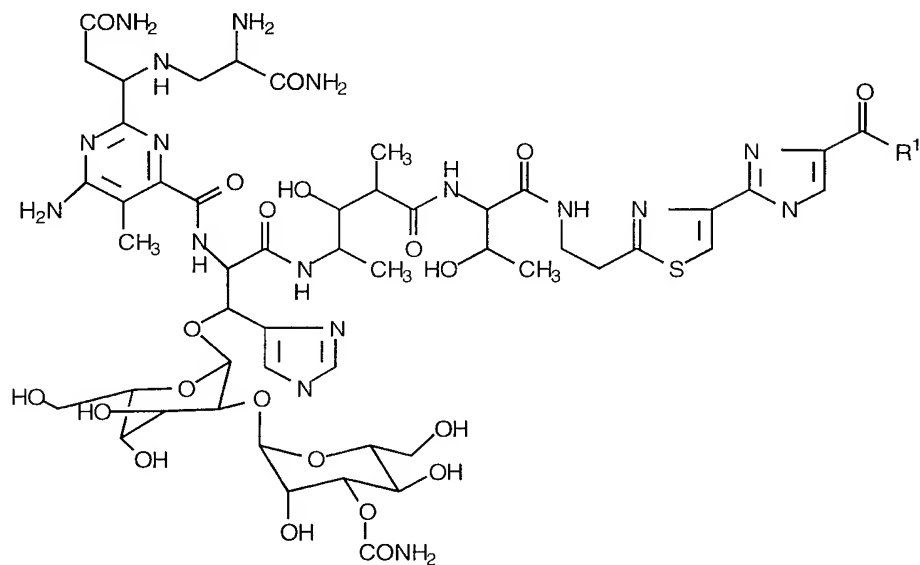
in which

R¹ is hydrogen or methyl;

R² is -NH₂, -OCH₃, -O(CH₂)₂OH, -NH(CH₂)₂SS(CH₂)₂NHAc, -NHCH-C≡CH, -NH(CH₂)₂SS(C₆H₄)NO₂, -O(CH₂)₂SS(CH₂)₂OH, -N=CH-NHOCH₃, -NH(C₆H₄)OH, -NH(CH₂)₂SS(CH₂)₂NHCO(CH₂)₂CH(NH₂)COOH



THE BLEOMYCIN GROUP OF FORMULA (2):

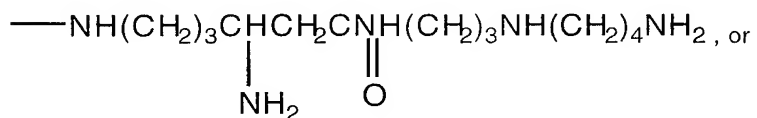
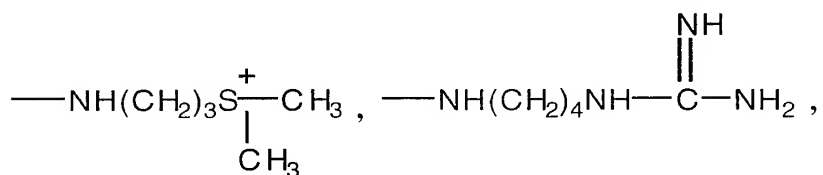


(2)

5 in which

R^1 is hydroxy, amino, C_1 - C_3 alkylamino.

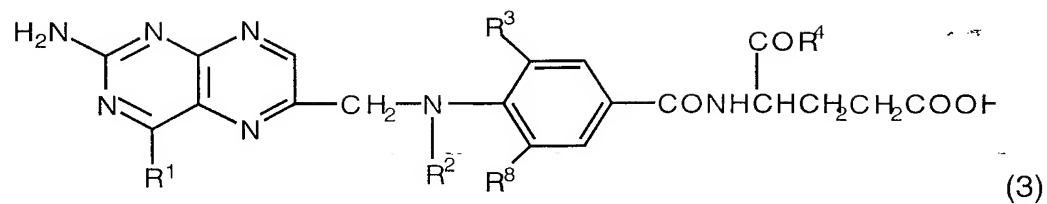
di(C_1 - C_3 alkyl)amino, C_4 - C_6 polymethylene amino,



10



THE METHOTREXATE GROUP OF FORMULA (3):



in which

R¹ is amino or hydroxy;

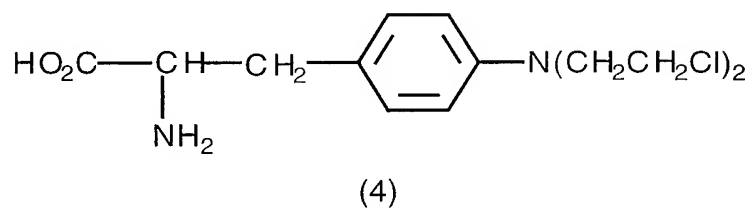
R² is hydrogen or methyl;

5 R³ is hydrogen, fluoro, chloro, bromo or iodo;

R⁴ is hydroxy or a moiety which complete a salt of the carboxylic acid.

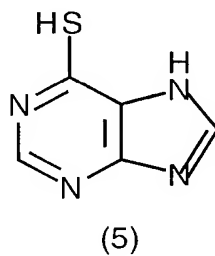
MELPHALAN OF FORMULA (4):

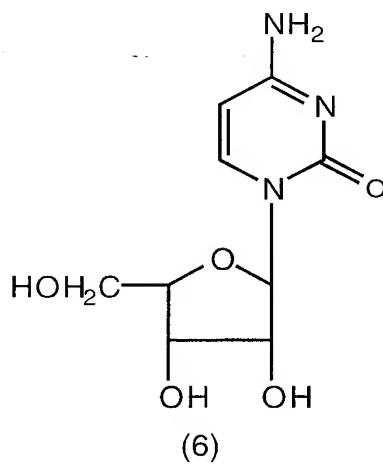
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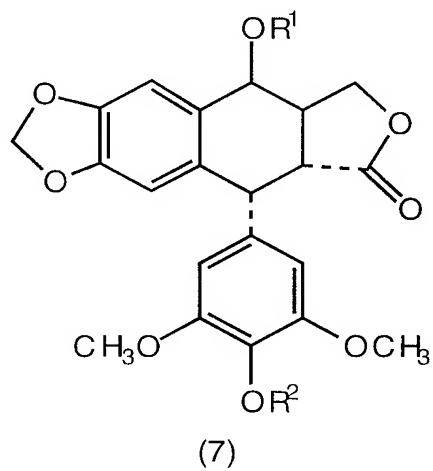
6-MERCAPTOPURINE OF FORMULA (5):

15



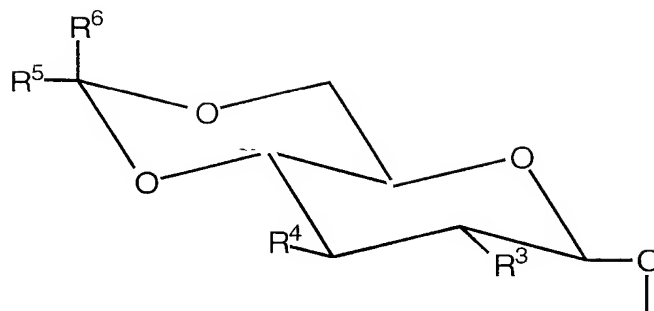
A CYTOSINE ARABINOSIDE OF FORMULA (6):

5

THE PODOPHYLLOTOXINS OF FORMULA (7):

10 wherein

R² is hydrogenR¹ is hydrogen or



wherein

R^3 is NH_2 , OH , OCH_3 , $\text{NH}(\text{C}_1\text{-C}_3 \text{ alkyl})$ or $\text{N}(\text{C}_1\text{-C}_3 \text{ alkyl})_2$

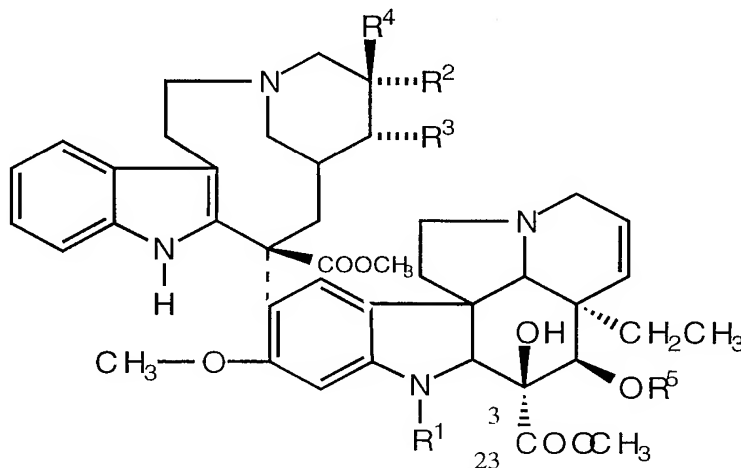
5 R^4 is OH , or NH_2 ,

R^5 is methyl or thienyl,

R^6 is hydrogen or methyl, or a phosphate salt thereof.

As used herein "(C₁-C₃ alkyl)" means a straight or branched carbon chain having from one to three carbon atoms; examples include methyl,
 10 ethyl, n-propyl and isopropyl.

THE VINCA ALKALOID GROUP OF DRUGS OF FORMULA (8):



15 in which

R^1 is H , CH_3 or CHO ;

when R^2 and R^3 are taken singly, R^3 is H , and one of R^4 and R^5 is ethyl and the other is H or OH :

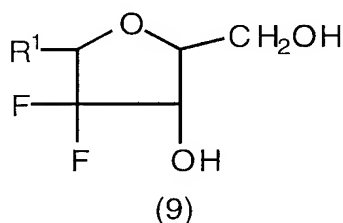
when R^2 and R^3 are taken together with the carbons to which they are attached, they form an oxirane ring in which case R^4 is ethyl;

R^5 is hydrogen, (C₁-C₃ alkyl)-CO, or chlorosubstituted (C₁-C₂ alkyl)-CO.

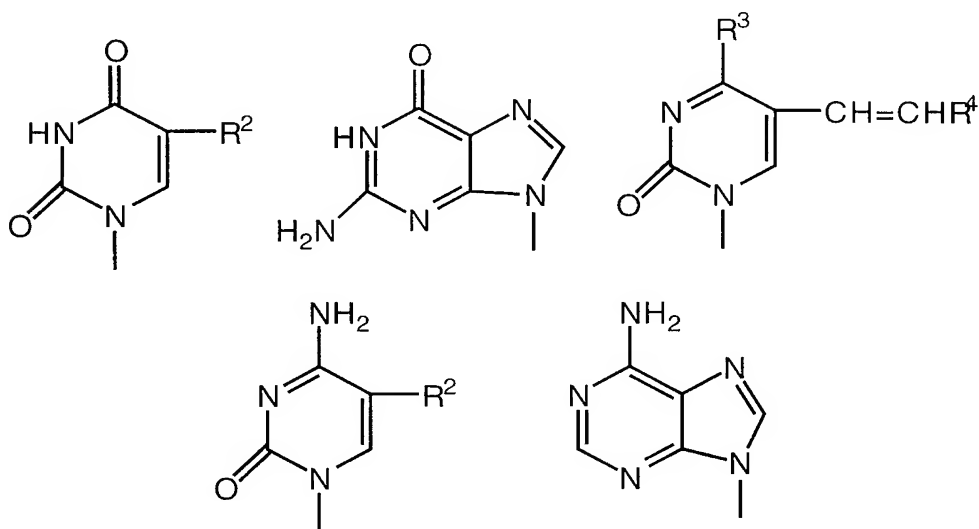
- 5 As used herein "C₁-C₃ alkyl" means a straight or branched carbon chain having from one to three carbon atoms; examples include methyl, ethyl, n-propyl and isopropyl.

DIFLUORONUCLEOSIDES OF FORMULA (9):

10



in which R^1 is a base of one of the formulae:



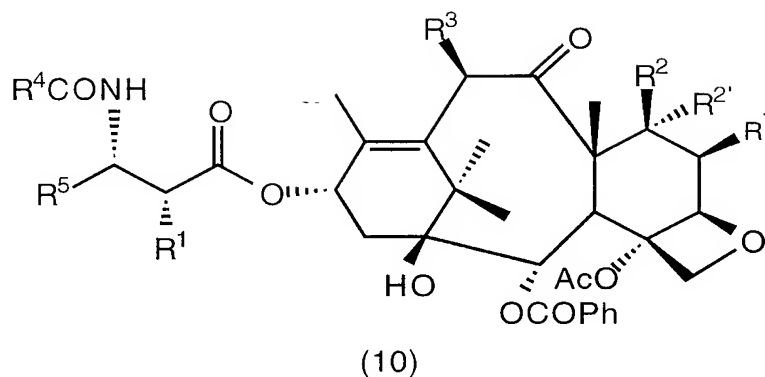
15

in which

R^2 is hydrogen, methyl, bromo, fluoro, chloro, or iodo;

R^3 is -OH or -NH₂;

R^4 is hydrogen, bromo, chloro, or iodo.

TAXOLS OF FORMULA (10):

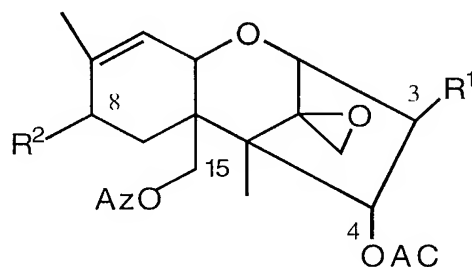
wherein

- 5 R¹ is hydroxy;
 R² is hydrogen or hydroxy;
 R^{2'} is hydrogen, hydroxy, or acetoxy;
 R⁷ is hydrogen or hydroxy;
 R³ is hydrogen, hydroxy, or acetoxy;
- 10 R⁴ is aryl, substituted aryl, C₁-6 alkyl, C₂-6 alkenyl, C₂-6 alkynyl or
 t-butoxy;
 R⁵ is C₁-6 alkyl, C₂-6 alkenyl, C₂-6 alkynyl, or -Z-R⁶;
 Z is a direct bond, C₁-6 alkyl or C₂-6 alkenyl;
 R⁶ is aryl, substituted aryl, C₃-6 cycloalkyl, thienyl or furyl.
- 15 As used herein, "alkyl" means a straight or branched saturated
 carbon chain having from one to six carbon atoms; examples include methyl,
 ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, isobutyl, t-butyl, n-pentyl, sec-
 butyl, isopentyl, and n-hexyl. "Alkenyl" means a straight or branched carbon
 chain having at least one carbon-carbon double bond, and having from two
 20 to six carbon atoms; examples include ethenyl, propenyl, isopropenyl,
 butenyl, isobutenyl, pentenyl, and hexenyl. "Alkynyl" means a straight or
 branched carbon chain having at least one carbon-carbon triple bond, and
 from two to six carbon atoms; examples include ethynyl, propynyl, butynyl,
 and hexynyl. "Aryl" means aromatic hydrocarbon having from six to ten
 25 carbon atoms; examples include phenyl and naphthyl. "Substituted aryl"
 means aryl substituted with at least one group selected from C₁-6

alkanoyloxy, hydroxy, halogen, C₁-6 alkyl, trifluoromethyl, C₁-6 alkoxy, aryl, C₂-6 alkenyl, C₁-6 alkanoyl, nitro, amino, and amido.

ANGUIDINES OF FORMULA (11):

5



(11)

wherein

R¹ is OH or O;

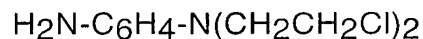
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R² is H or O;

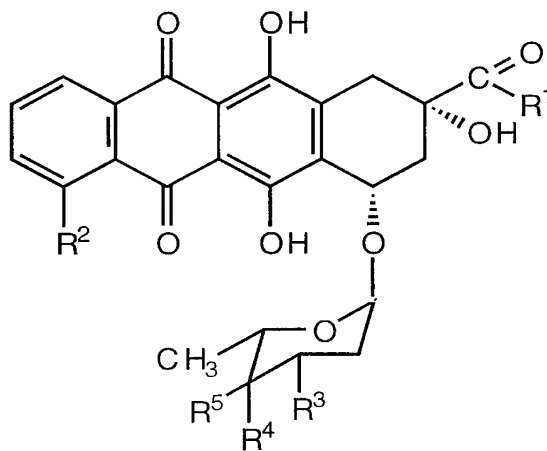
Anguidine can be targeted at the C-3, C-4, C-8 or C-15 positions, as an ester or hydrazone

PHENYLENE DIAMINE MUSTARD (12):

15



THE ANTHRACYCLINES ANTIBIOTICS OF FORMULA (13):



(13)

20

wherein

R¹ is -CH₃, -CH₂OH, -CH₂OCO(CH₂)₃CH₃ or -CH₂OCOCH(OC₂H₅)₂

R² is -OCH₃, -OH or -H

5 R³ is -NH₂, -NHCOCF₃, 4-morpholinyl, 3-cyano-4-morpholinyl, 1-piperidinyl, 4-methoxy-1-piperidinyl, benzylamine, dibenzylamine, cyanomethylamine, 1-cyano-2-methoxyethyl amine, or NH-(CH₂)₄-CH(OAc)₂;

R⁴ is -OH, -OTHP, or -H; and

10 R⁵ is -OH or -H provided that R⁵ is not -OH when R⁴ is -OH or -OTHP.

One skilled in the art understands that structural Formula (13) includes compounds which are drugs, or are derivatives of drugs, which
15 have acquired in the art different generic or trivial names. Table I, which follows, represents a number of anthracycline drugs and their generic or trivial names and which are especially preferred for use in the present invention.

Of the compounds shown in Table I, the most highly preferred drug
20 is Doxorubicin. Doxorubicin (also referred to herein as "DOX") is that anthracyclin shown in the formula of Table I in which R¹ is -CH₂OH, R² is -OCH₃, R³ is -NH₂, R⁴ is -OH, and R⁵ is -H.

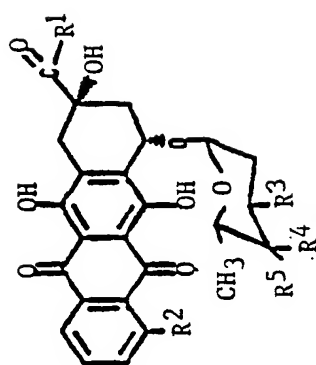


Table I

Compound	R1	R2	R3	R4	R5
Daunorubicin*	CH ₂ OH	OCH ₃	NH ₂	OH	H
Doxorubicin	CH ₂ OH	OCH ₃	NH ₂	OH	H
Detorubicin	CH ₂ OCOCH(OC ₂ H ₅) ₂	OCH ₃	NH ₂	OH	H
Carminomycin	CH ₃	OH	NH ₂	OH	H
Idarubicin	CH ₃	H	NH ₂	OH	H
Epilubicin	CH ₂ OH	OCH ₃	NH ₂	H	OH
Esorubicin	CH ₂ OH	OCH ₃	NH ₂	H	H
THP	CH ₂ OH	OCH ₃	NHCOCH ₃	OTHP	H
AD-32	CH ₂ OCO(CH ₂) ₃ CH ₃	OCH ₃	NHCOCH ₃	OH	H
Morpholino-Dox	CH ₂ OH	OCH ₃		OH	H
Cyano-morpholino-Dox	CH ₂ OH	OCH ₃		OH	H
DAPDox	CH ₂ OH	OCH ₃	-NH(CH ₂) ₄ CH(OAc) ₂	OH	H

*"Daunomycin" is an alternate name for daunorubicin

The most highly preferred drugs are taxol, mitomycin C, and the anthracycline antibiotic agents of Formula (13), described previously.

The Ligand

5 The "ligand" includes within its scope any molecule that specifically binds or reactively associates or complexes with a receptor or other receptive moiety associated with a given target cell population. This cell reactive molecule, to which the drug reagent is linked via the linker in the conjugate, can be any molecule that binds to, complexes with or reacts with
10 the cell population sought to be therapeutically or otherwise biologically modified and, which possesses a free reactive sulfhydryl (-SH) group or can be modified to contain such a sulfhydryl group. The cell reactive molecule acts to deliver the therapeutically active drug moiety to the particular target cell population with which the ligand reacts. Such molecules include, but
15 are not limited to, large molecular weight proteins such as, for example, antibodies, smaller molecular weight proteins, polypeptides or peptide ligands, and non-peptidyl ligands.

 The non-immunoreactive protein, polypeptide, or peptide ligands which can be used to form the conjugates of this invention may include, but
20 are not limited to, transferrin, epidermal growth factors ("EGF"), bombesin, gastrin, gastrin-releasing peptide, platelet-derived growth factor, IL-2, IL-6, tumor growth factors ("TGF"), such as TGF- α and TGF- β , vaccinia growth factor ("VGF"), insulin and insulin-like growth factors I and II. Non-peptidyl ligands may include, for example, carbohydrates, lectins, and apoprotein
25 from low density lipoprotein.

 The immunoreactive ligands comprise in antigen-recognizing immunoglobulin (also referred to as "antibody"), or an antigen-recognizing fragment thereof. Particularly preferred immunoglobulins are those immunoglobulins which can recognize a tumor-associated antigen. As
30 used, "immunoglobulin" may refer to any recognized class or subclass of immunoglobulins such as IgG, IgA, IgM, IgD, or IgE. Preferred are those immunoglobulins which fall within the IgG class of immunoglobulins. The immunoglobulin can be derived from any species. Preferably, however, the

immunoglobulin is of human, murine, or rabbit origin. Furthermore, the immunoglobulin may be polyclonal or monoclonal, preferably monoclonal.

As noted, one skilled in the art will appreciate that the invention also encompasses the use of antigen recognizing immunoglobulin fragments. Such immunoglobulin fragments may include, for example, the Fab', F(ab')₂, F_V or Fab fragments, or other antigen recognizing immunoglobulin fragments. Such immunoglobulin fragments can be prepared, for example, by proteolytic enzyme digestion, for example, by pepsin or papain digestion, reductive alkylation, or recombinant techniques.

The materials and methods for preparing such immunoglobulin fragments are well-known to those skilled in the art. See generally, Parham, J. Immunology, 131, 2895 (1983); Lamoyi et al., J. Immunological Methods, 56, 235 (1983); Parham, id., 53, 133 (1982); and Matthew et al., id., 50, 239 (1982).

The immunoglobulin can be a "chimeric antibody" as that term is recognized in the art. Also the immunoglobulin may be a "bifunctional" or "hybrid" antibody, that is, an antibody which may have one arm having a specificity for one antigenic site. such as a tumor associated antigen while the other arm recognizes a different target, for example, a hapten which is, or to which is bound, an agent lethal to the antigen-bearing tumor cell.

Alternatively, the bifunctional antibody may be one in which each arm has specificity for a different epitope of a tumor associated antigen of the cell to be therapeutically or biologically modified. In any case, the hybrid antibodies have a dual specificity, preferably with one or more binding sites specific for the hapten of choice or more or more binding sites specific for a target antigen, for example, an antigen associated with a tumor, an infectious organism, or other disease state.

Biological bifunctional antibodies are described, for example, in European Patent Publication, EPA 0 105 360, to which those skilled in the art are referred. Such hybrid or bifunctional antibodies may be derived, as noted, either biologically, by cell fusion techniques, or chemically, especially with cross-linking agents or disulfide bridge-forming reagents, and may be comprised of whole antibodies and/or fragments thereof. Methods for obtaining such hybrid antibodies are disclosed, for example, in PCT

Application WO83/03679, published October 27, 1983, and published
European Application EPA 0 217 577, published April 8, 1987, both of
which are incorporated herein by reference. Particularly preferred
bifunctional antibodies are those biologically prepared from a "polydoma" or
5 "quadroma" or which are synthetically prepared with cross-linking agents
such as bis-(maleimido)-methyl ether ("BMME"), or with other cross-linking
agents familiar to those skilled in the art.

In addition the immunoglobulin may be a single chain antibody
("SCA"). These may consist of single chain Fv fragments ("scFv") in which
10 the variable light ("VL") and variable heavy ("VH") domains are linked by a
peptide bridge or by disulfide bonds. Also, the immunoglobulin may consist
of single VH domains (dAbs) which possess antigen-binding activity. See,
e.g., G. Winter and C. Milstein, Nature, 349, 295 (1991); R. Glockshuber et
al., Biochemistry 29, 1362 (1990); and E. S. Ward et al., Nature 341, 544
15 (1989).

Especially preferred for use in the present invention are chimeric
monoclonal antibodies, preferably those chimeric antibodies having
specificity toward a tumor associated antigen. As used herein, the term
"chimeric antibody" refers to a monoclonal antibody comprising a variable
20 region, i.e., binding region, from one source or species and at least a portion
of a constant region derived from a difference source of species, usually
prepared by recombinant DNA techniques. Chimeric antibodies comprising
a murine variable region and a human constant region are especially
preferred in certain applications of the invention, particularly human therapy,
25 because such antibodies are readily prepared and may be less
immunogenic than purely murine monoclonal antibodies. Such
murine/human chimeric antibodies are the product of expressed
immunoglobulin genes comprising DNA segments encoding murine
immunoglobulin constant regions. Other forms of chimeric antibodies
30 encompassed by the invention are those in which the class or subclass has
been modified or changed from that of the original antibody. Such "chimeric"
antibodies are also referred to as "class-switched antibodies". Methods for
producing chimeric antibodies involve conventional recombinant DNA and

gene transfection techniques now well known in the art. See, e.g., Morrison, S. L., et al., Proc. Nat'l Acad. Sci, 81 6851 (1984).

Encompassed by the term "chimeric antibody" is the concept of "humanized antibody", that is those antibodies in which the framework or
5 "complementarity determining regions ("CDR") have been modified to comprise the CDR of an immunoglobulin of different specificity as compared to that of the parent immunoglobulin. In a preferred embodiment, a murine CDR is grafted into the framework region of a human antibody to prepare the "humanized antibody". See, e.g., L. Riechmann et al., Nature 332, 323
10 (1988); M. S. Neuberger et al., Nature 314, 268 (1985). Particularly preferred CDR's correspond to those representing sequences recognizing the antigens noted above for the chimeric and bifunctional antibodies. The reader is referred to the teaching of EPA 0 239 400 (published September 30, 1987), incorporated herein by reference, for its teaching of CDR modified
15 antibodies.

One skilled in the art will recognize that a bifunctional-chimeric antibody can be prepared which would have the benefits of lower immunogenicity of the chimeric or humanized antibody, as well as the flexibility, especially for therapeutic treatment, of the bifunctional antibodies
20 described above. Such bifunctional-chimeric antibodies can be synthesized, for instance, by chemical synthesis using cross-linking agents and/or recombinant methods of the type described above. In any event, the present invention should not be construed as limited in scope by any particular method of production of an antibody whether bifunctional,
25 chimeric, bifunctional-chimeric, humanized, or an antigen-recognizing fragment or derivative thereof.

In addition, the invention encompasses within its scope immunoglobulins (as defined above) or immunoglobulin fragments to which are fused active proteins, for example, an enzyme of the type disclosed in
30 Neuberger, et al., PCT application, WO86/01533, published March 13, 1986. The disclosure of such products is incorporated herein by reference.

As noted, "bifunctional", "fused", "chimeric" (including humanized), and "bifunctional-chimeric" (including humanized) antibody constructions also include, within their individual contexts constructions comprising

antigen recognizing fragments. As one skilled in the art will recognize, such fragments could be prepared by traditional enzymatic cleavage of intact bifunctional, chimeric, humanized, or chimeric-bifunctional antibodies. If, however, intact antibodies are not susceptible to such cleavage, because of the nature of the construction involved, the noted constructions can be prepared with immunoglobulin fragments used as the starting materials; or, if recombinant techniques are used, the DNA sequences, themselves, can be tailored to encode the desired "fragment" which, when expressed, can be combined in vivo or in vitro, by chemical or biological means, to prepare the final desired intact immunoglobulin "fragment". It is in this context, therefore, that the term "fragment" is used.

Furthermore, as noted above, the immunoglobulin (antibody), or fragment thereof, used in the present invention may be polyclonal or monoclonal in nature. Monoclonal antibodies are the preferred immunoglobulins, however. The preparation of such polyclonal or monoclonal antibodies now is well known to those skilled in the art who, of course, are fully capable of producing useful immunoglobulins which can be used in the invention. See, e.g., G. Kohler and C. Milstein, Nature 256, 495 (1975). In addition, hybridomas and/or monoclonal antibodies which are produced by such hybridomas and which are useful in the practice of the present invention are publicly available from sources such as the American Type Culture Collection ("ATCC") 12301 Parklawn Drive, Rockville, Maryland 20852 or, commercially, for example, from Boehringer-Mannheim Biochemicals, P.O. Box 50816, Indianapolis, Indiana 46250.

Particularly preferred monoclonal antibodies for use in the present invention are those which recognize tumor associated antigens. Such monoclonal antibodies, are not to be so limited, however, and may include, for example, the following:

<u>Antigen Site Recognized</u>	<u>Monoclonal Antibodies</u>	<u>Reference</u>
Lung Tumors	KS1/4	N. M. Varki <u>et al.</u> , <u>Cancer Res.</u> 44:681, 1984.
	534,F8;604A9	F. Cuttitta <u>et al.</u> , in: G. L. Wright (ed) <u>Monoclonal Antibodies and Cancer</u> , Marcel Dekker, Inc., NY., p. 161, 1984.
Squamous Lung	G1, LuCa2, LuCa3, LuCa4	Kyoizumi <u>et al.</u> , <u>Cancer Res.</u> , 45:3274, 1985.
Small Cell Lung Cancer	TFS-2	Okabe <u>et al.</u> , <u>Cancer Res.</u> , 45:1930, 1985.
Colon Cancer	11.285.14 14.95.55	G. Rowland <u>et al.</u> , <u>Cancer Immunol. Immunother.</u> , 19:1, 1985.
	NS-3a-22,NS-10 NS-19-9,NS-33a NS-52a,17-1A	Z. Steplewski <u>et al.</u> , <u>Cancer Res.</u> , 41:2723, 1981.
Carcinoembryon ic	MoAb 35 or ZCE025	Acolla, R. S. <u>et al.</u> , <u>Proc. Nat. Acad. Sci., (USA)</u> , 77:563, 1980.
Melanoma	9.2.27	T. F. Bumol and R. A. Reiseld, <u>Proc. Natl. Acad. Sci., (USA)</u> , 79:1245, 1982.

p97	96.5	K. E. Hellstrom <u>et al.</u> , <u>Monoclonal Antibodies and</u> <u>Cancer, loc. cit.</u> p. 31.
Antigen T65	T101	Boehringer-Mannheim P.O. Box 50816 Indianapolis, IN 46250
Ferritin	Antiferrin	Boehringer-Mannheim P.O. Box 50816 Indianapolis, IN 46250
	R24	W. G. Dippold <u>et al.</u> , <u>Proc.</u> <u>Natl. Acad. Sci. (USA)</u> , 77:6114, 1980.
Neuroblastoma	P1 153/3	R. H. Kennet and F. Gilbert, <u>Science</u> , 203:1120, 1979.
	MIN 1	J. T. Kemshead in <u>Monoclonal Antibodies and</u> <u>Cancer, loc. cit.</u> p. 49.
	UJ13A	Goldman <u>et al.</u> , <u>Pediatrics</u> , 105:252, 1984.
Glioma	BF7, GE2, CG12	N. de Tribolet <u>et al.</u> , in <u>Monoclonal Antibodies and</u> <u>Cancer, loc. cit.</u> p. 81.

Ganglioside	L6	I. Hellstrom <u>et al.</u> , <u>Proc. Natl. Acad. Sci. (USA)</u> , 83:7059 (1986); U.S. Patent Nos. 4,906,562, issued March 6, 1990 and 4,935,495, issued June 19, 1990.
	Chimeric L6	U.S. Serial No. 07/923,244, filed October 27, 1986, equivalent to PCT Patent Publication, WO 88/03145, published May 5, 1988.
Lewis Y	BR64	U.S. Serial No. 07/289,635, filed December 22, 1988, and U.S. Serial No. 07/443,696, filed November 29, 1989, equivalent to European Patent Publication, EP A 0 375 562, published June 27, 1990.
fucosylated Lewis Y	BR96, Chimeric BR96	U.S. Serial No. 07/374,947, filed June 30, 1989, and U.S. Serial No. 07/544,246, filed June 26, 1990, equivalent to PCT Patent Publication, WO 91/00295, published January 10, 1991.

Breast Cancer	B6.2, B72.3	D. Colcher <u>et al.</u> , in <u>Monoclonal Antibodies and Cancer, loc. cit.</u> p. 121.
Osteogenic Sarcoma	791T/48, 791T/36	M. J. Embleton, <u>ibid</u> , p. 181.
Leukemia	CALL 2 anti-idiotypic	C. T. Teng <u>et al.</u> , <u>Lancet</u> , 1:01, 1982. R.A. Miller <u>et al.</u> , <u>N. Engl. J. Med.</u> , 306:517, 1982.
Ovarian Cancer	OC 125	R. C. Bast <u>et al.</u> , <u>J. Clin. Invest.</u> , 68:1331, 1981.
Prostrate Cancer	D83.21, P6.2, Turp-27	J. J. Starling <u>et al.</u> , in <u>Monoclonal Antibodies and Cancer, loc. cit.</u> p. 253.
Renal Cancer	A6H, D5D	P. H. Lange <u>et al.</u> , <u>Surgery</u> , 98:143, 1985.

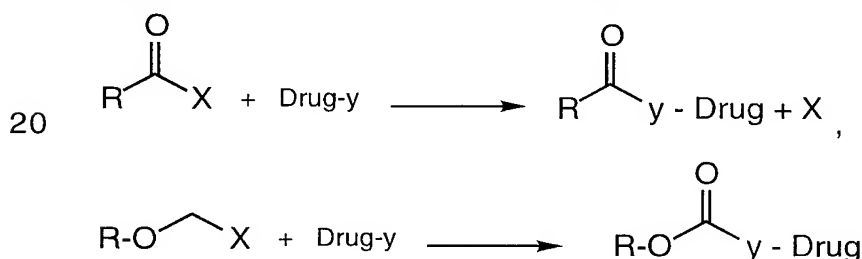
- In the most preferred embodiment, the ligand containing conjugate is derived from chimeric antibody BR96, "ChiBR96", disclosed in U.S. Serial No. 07/544,246, filed June 26, 1990, and which is equivalent to PCT
- 5 Published Application, WO 91/00295, published January 10, 1991.
- ChiBR96 is an internalizing murine/human chimeric antibody and is reactive, as noted, with the fucosylated Lewis Y antigen expressed by human carcinoma cells such as those derived from breast, lung, colon, and ovarian carcinomas. The hybridoma expressing chimeric BR96 and identified as
- 10 ChiBR96 was deposited on May 23, 1990, under the terms of the Budapest Treaty, with the American Type Culture Collection ("ATCC"), 12301 Parklawn Drive, Rockville, Maryland 20852. Samples of this hybridoma are available

under the accession number ATCC 10460. ChiBR96 is derived, in part, from its source parent, BR96. The hybridoma expressing BR96 was deposited, on February 21, 1989, at the ATCC, under the terms of the Budapest Treaty and is available under the accession number HB 10036. The desired hybridoma is cultured and the resulting antibodies are isolated from the cell culture supernatant using standard techniques now well known in the art. See, e.g., "Monoclonal Hybridoma Antibodies: Techniques and Applications", Hurell (ed.) (CRC Press, 1982).

Thus, as used "immunoglobulin" or "antibody" encompasses within its meaning all of the immunoglobulin/antibody forms or constructions noted above.

Synthesis of the Compounds of the Invention

The linkers, drug/linkers, and conjugates of the invention can be made techniques taught herein, known in the art, or can be made via routine experimentation using as guidance the techniques taught herein and/or known in the art. The attachment of the drug to the linker is accomplished by reacting a nucleophilic atom of the drug (O, N or S) to an electrophilic atom (C, S, P) on either the self-immolating spacer or the carboxy terminus of the peptide. This is illustrated as follows:

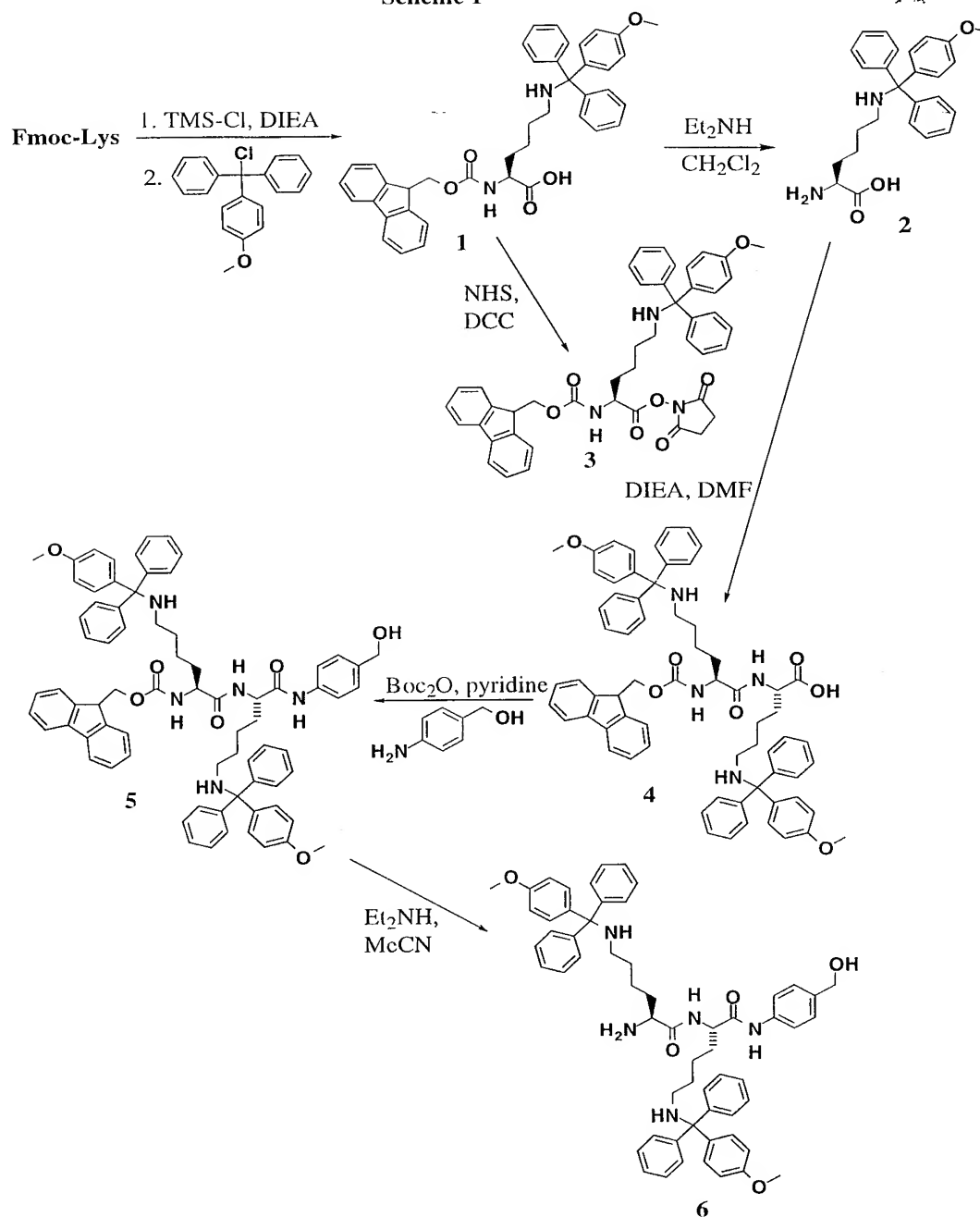


where: X = leaving group such as Cl⁻, Br⁻, tosylate, N-hydroxysuccinimide
 y = nucleophilic group such as OH, NH₂, SH, NH-lower alkyl
 R = rest of linker.

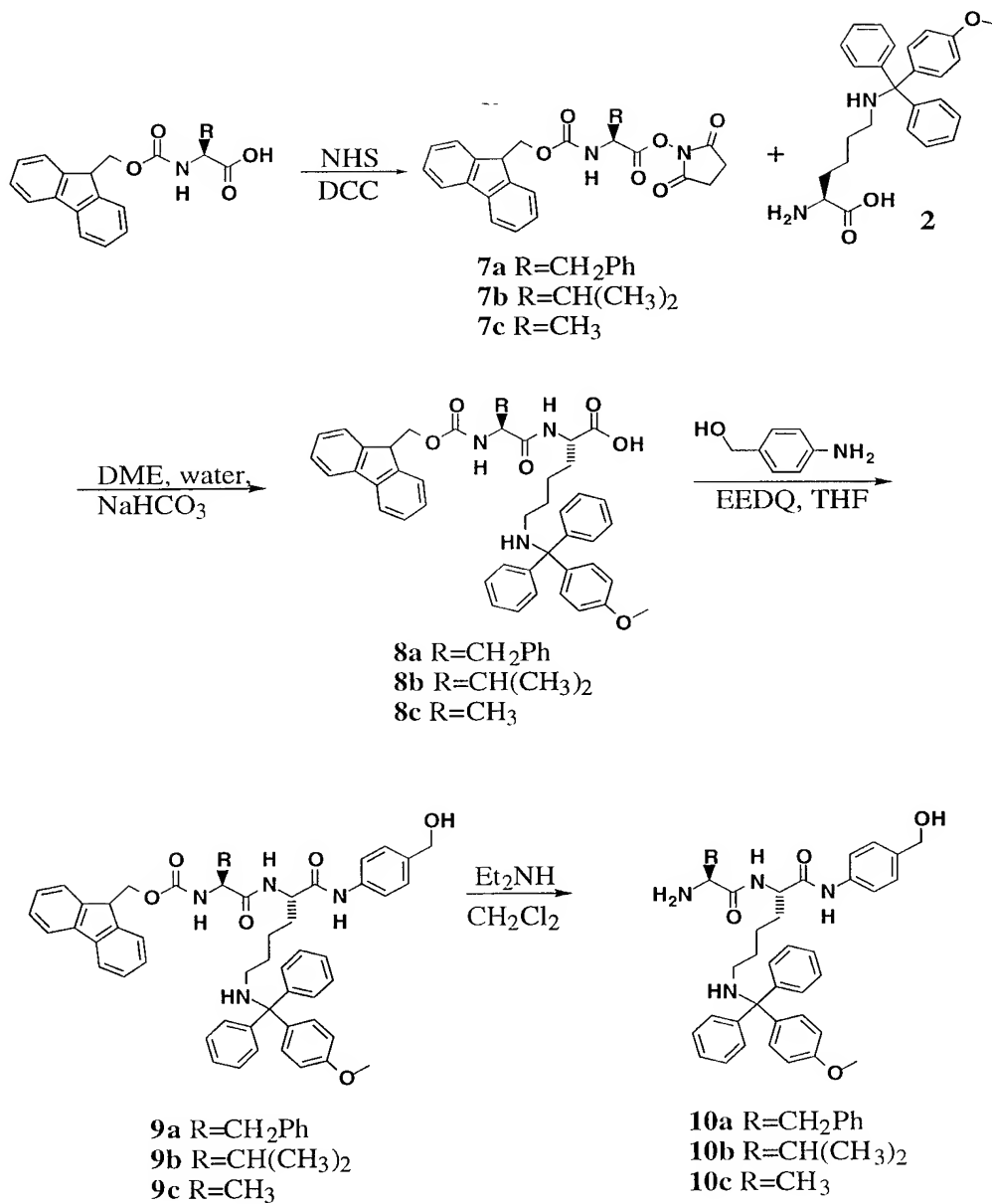
Linkage to an aldehyde or ketone can be effected by having the aldehyde or ketone in the form of an enol or N-loweralkyl enamine. It is expected that on release of the free enamine, it will spontaneously hydrolyze to an aldehyde or ketone.

The following reaction schemes are illustrative preparative techniques (compound numbers correspond to Example numbers):

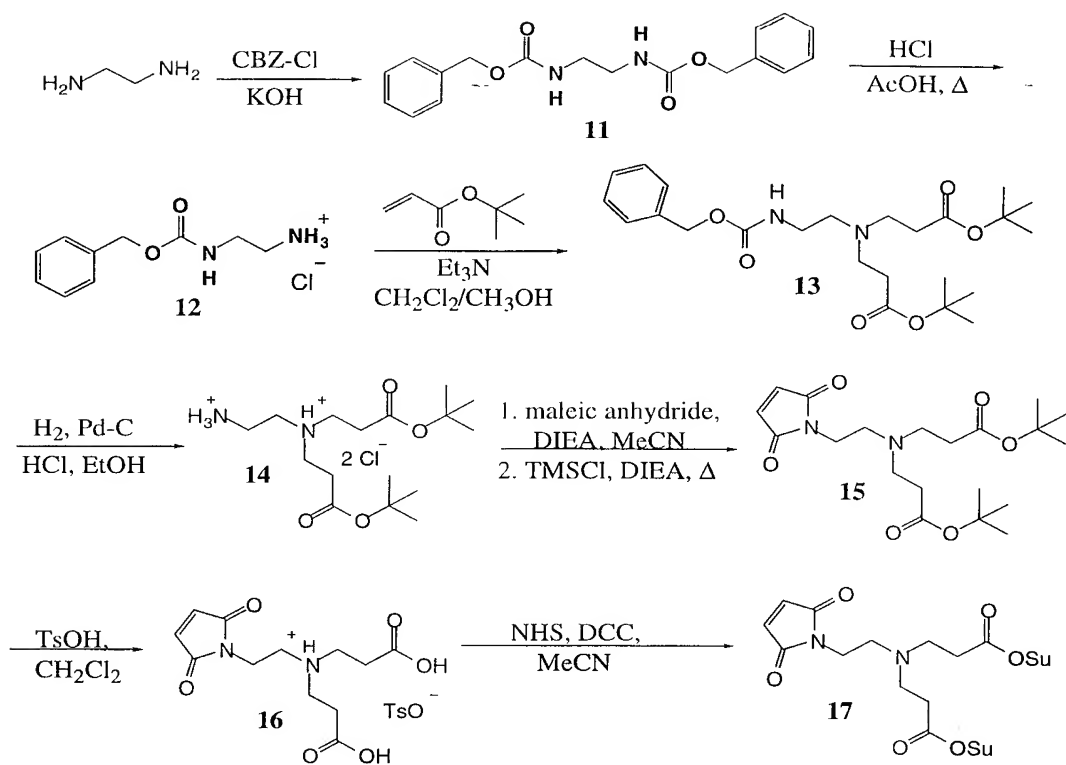
Scheme 1



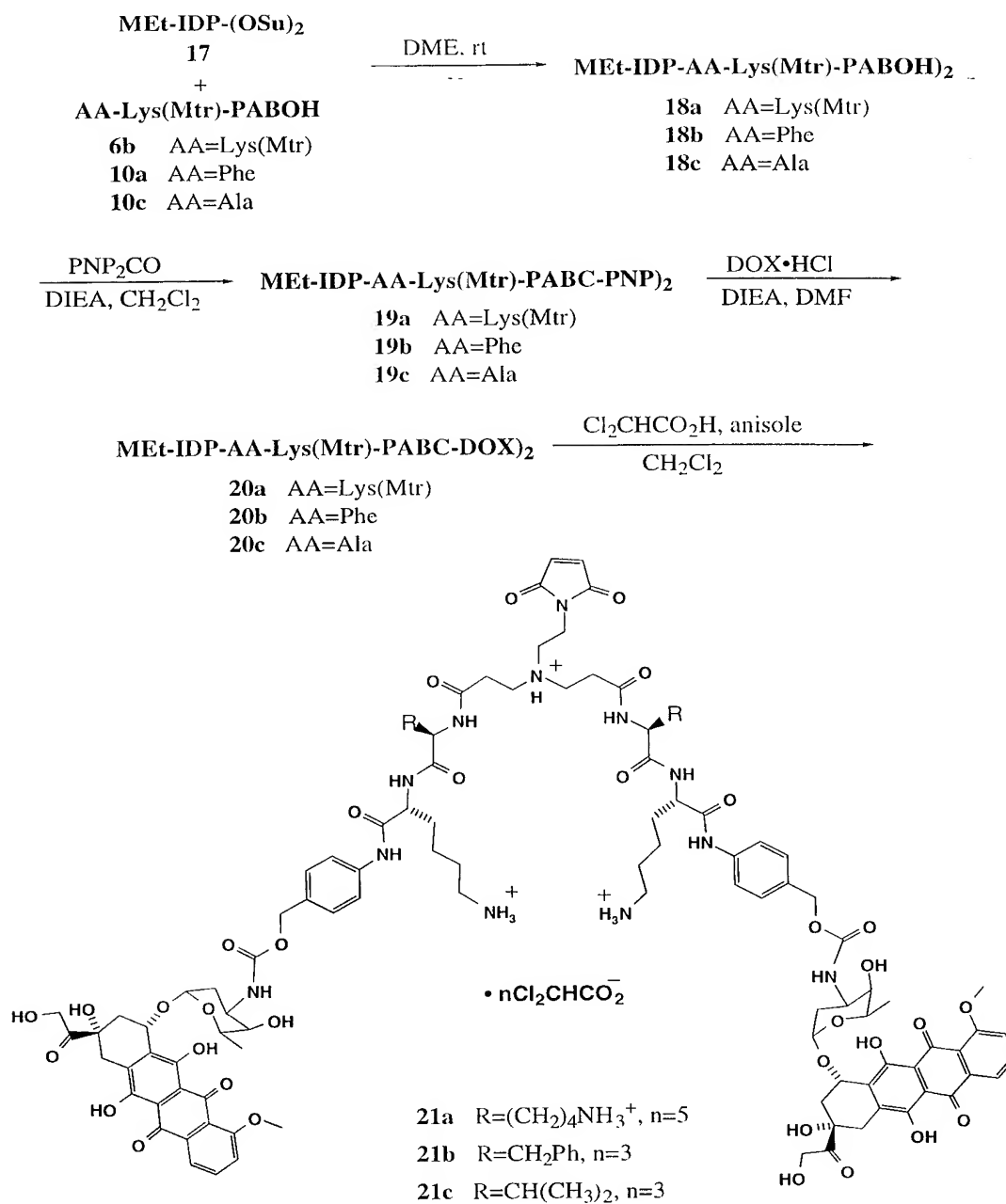
Scheme 2



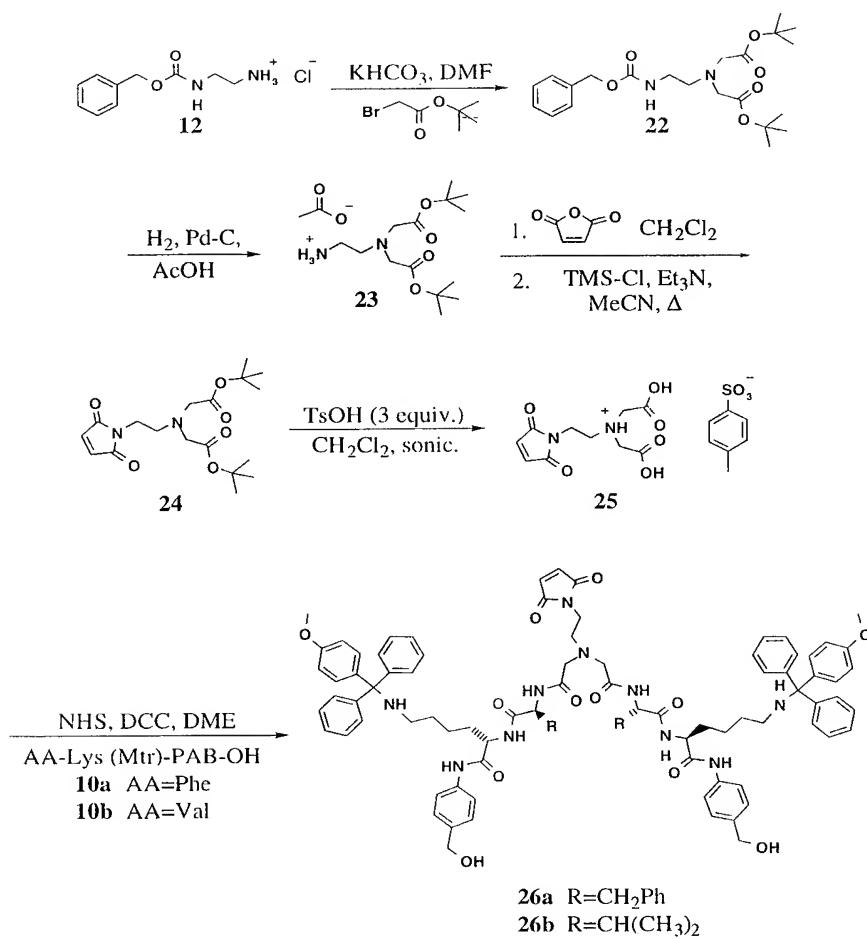
Scheme 3 (Part I)



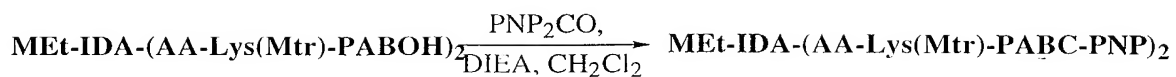
Scheme 3 (Part II)



Scheme 4 (Part I)



Scheme 4 (Part II)

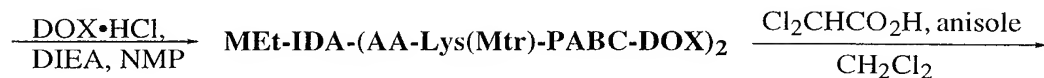


26a AA=Phe

26b AA=Val

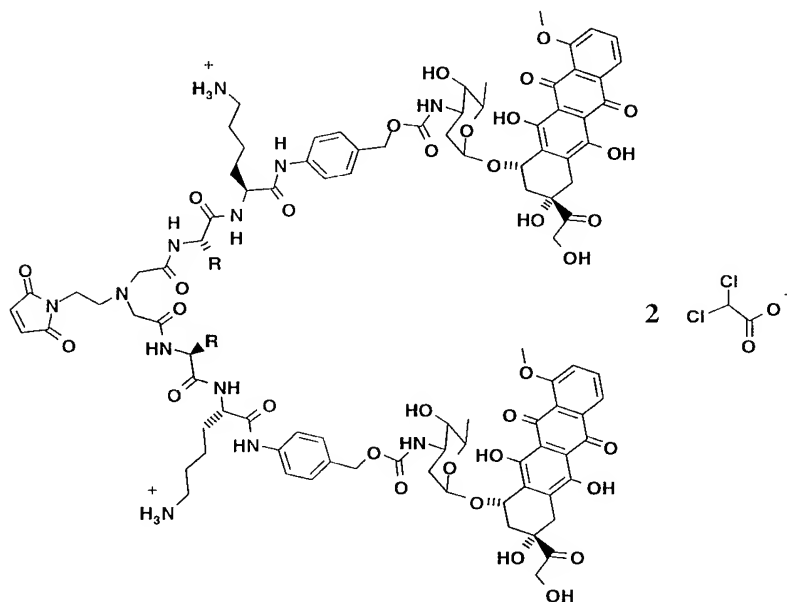
27a AA=Phe

27b AA=Val

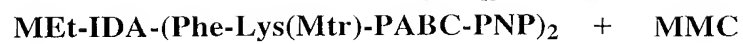


28a AA=Phe

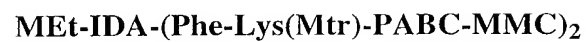
28b AA=Val

29a R=CH₂Ph29b R=CH(CH₃)₂

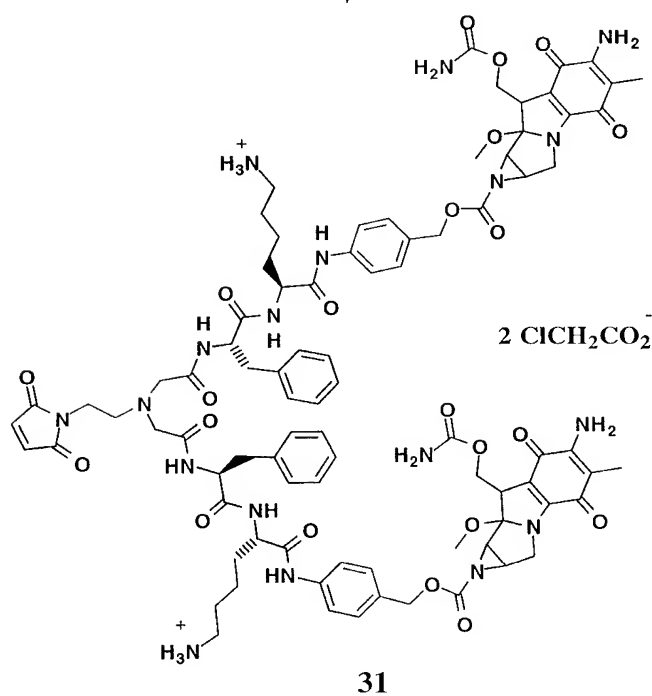
Scheme 5

**27a**

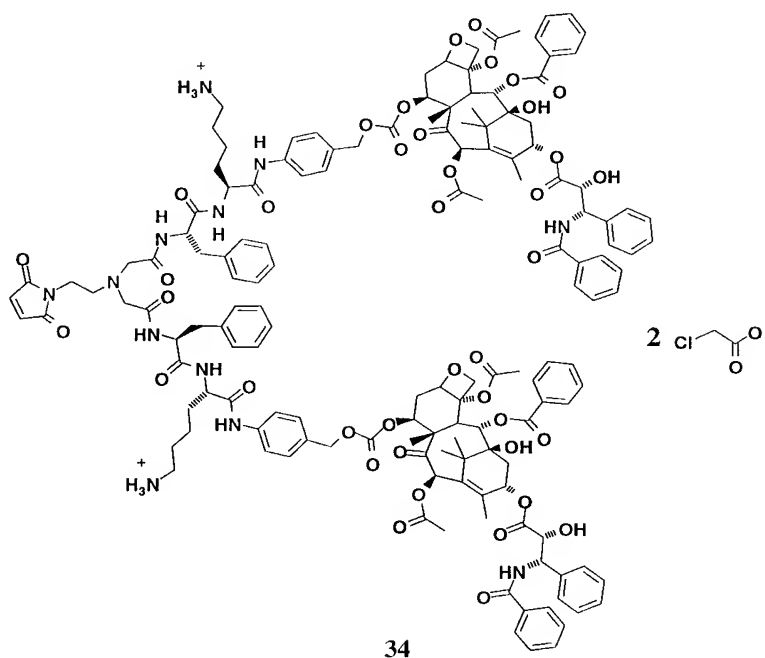
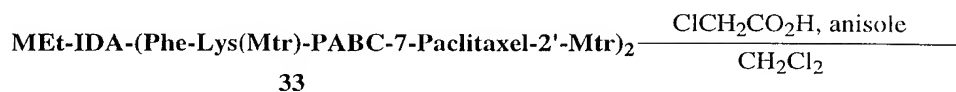
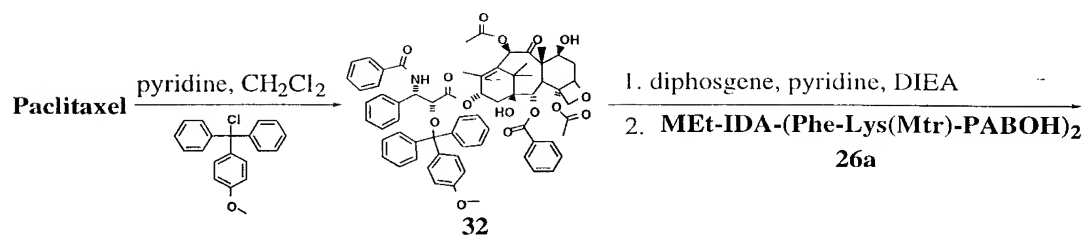
HOBT, DIEA
NMP, sieves

**30**

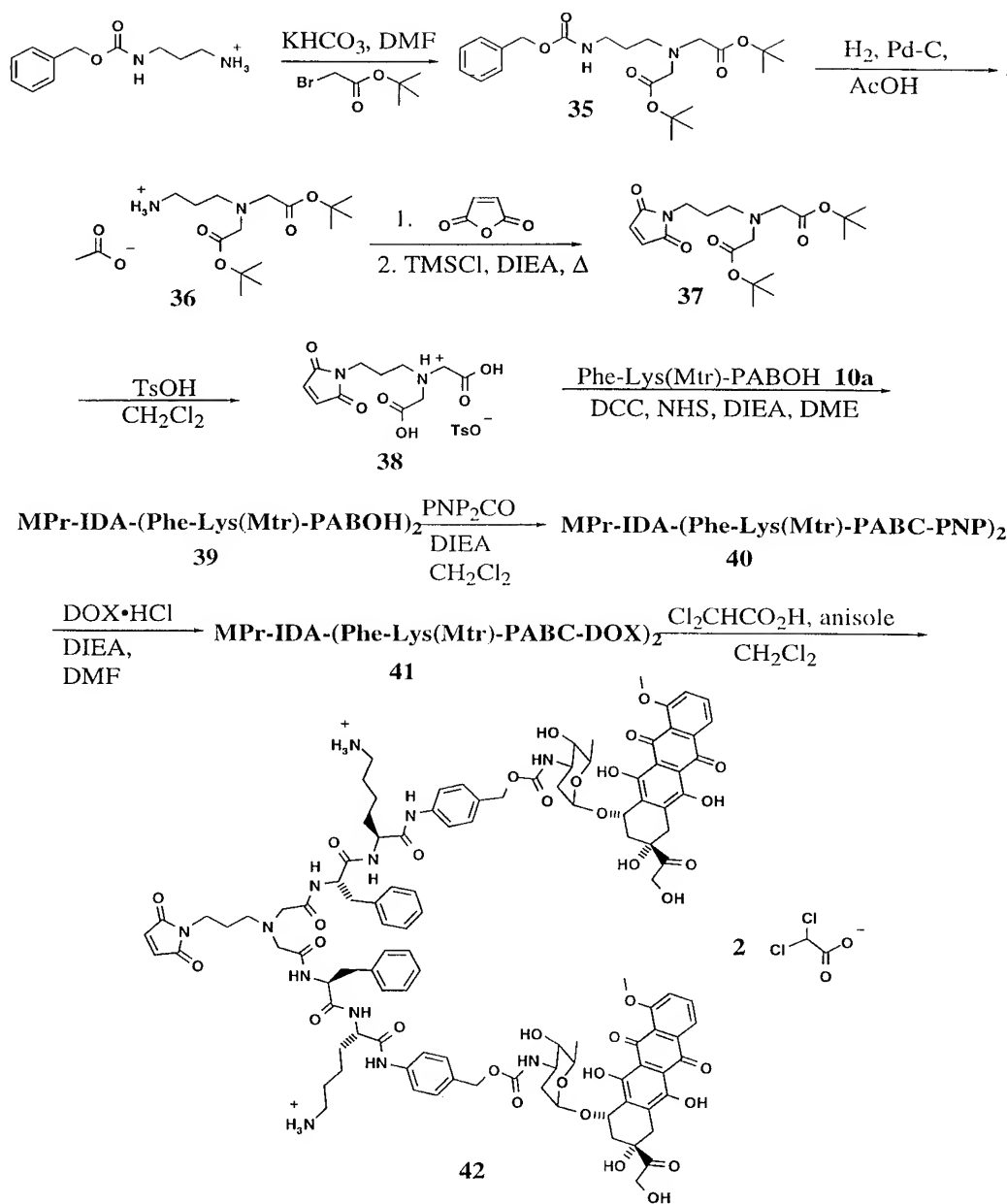
$\text{ClCH}_2\text{CO}_2\text{H}$,
anisole, CH_2Cl_2



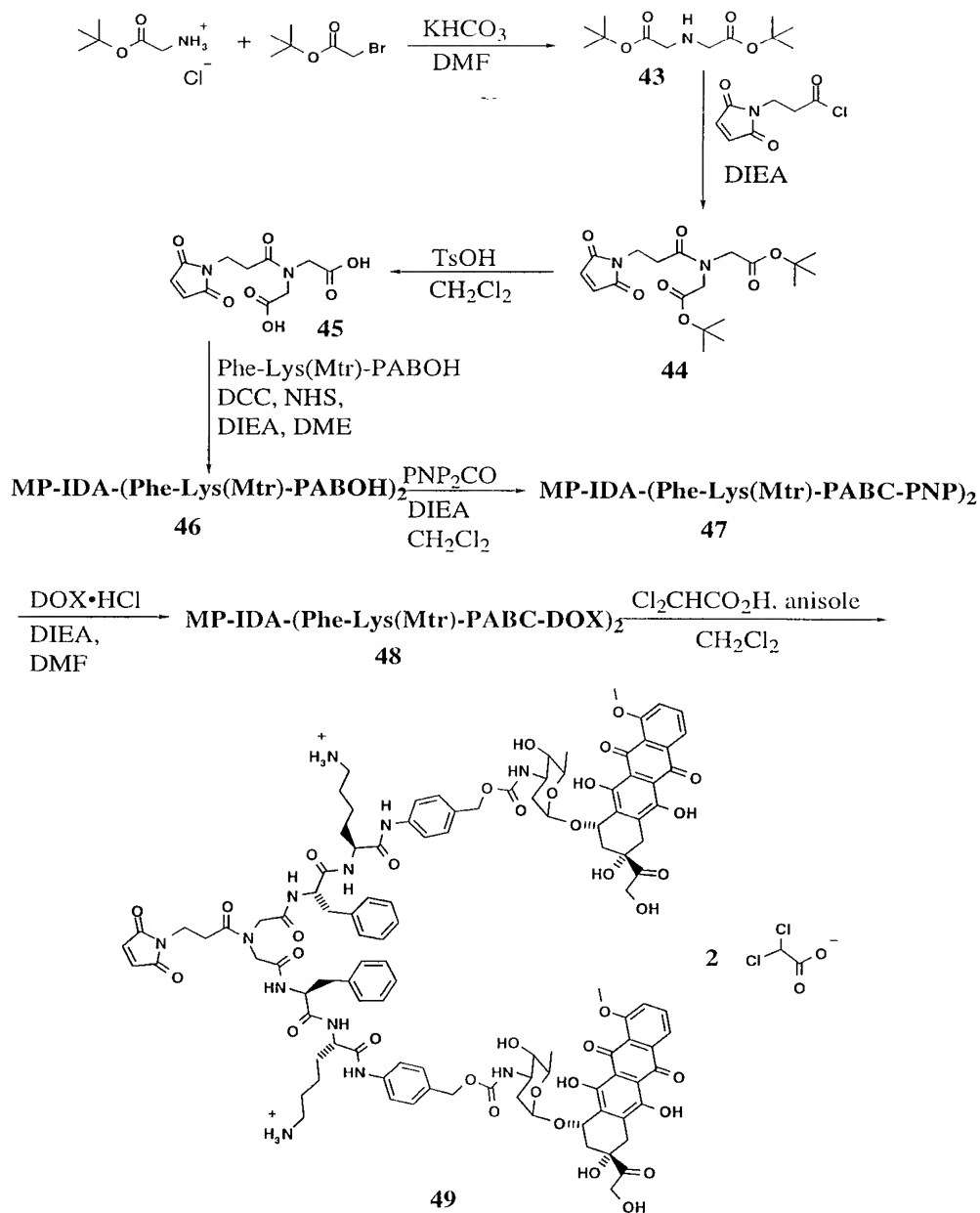
Scheme 6



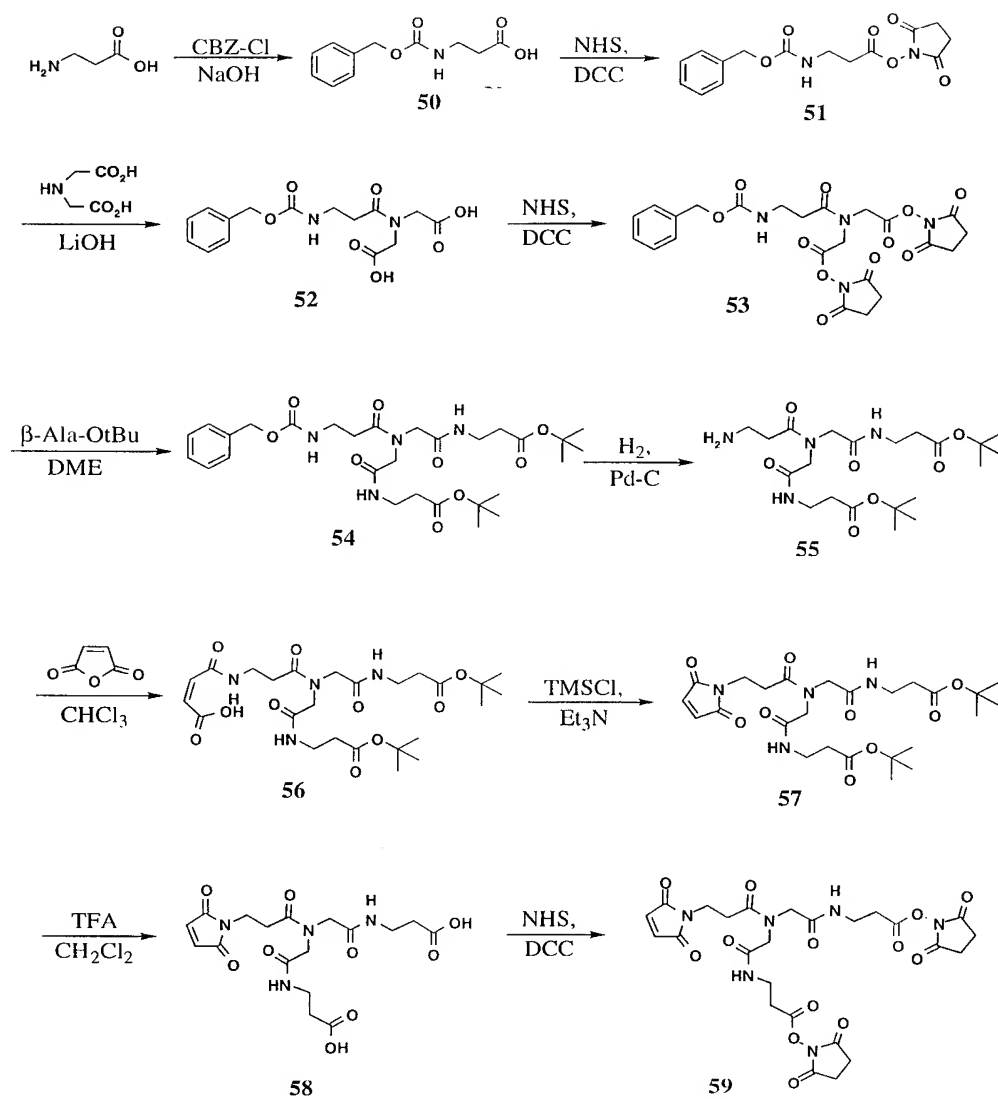
Scheme 7



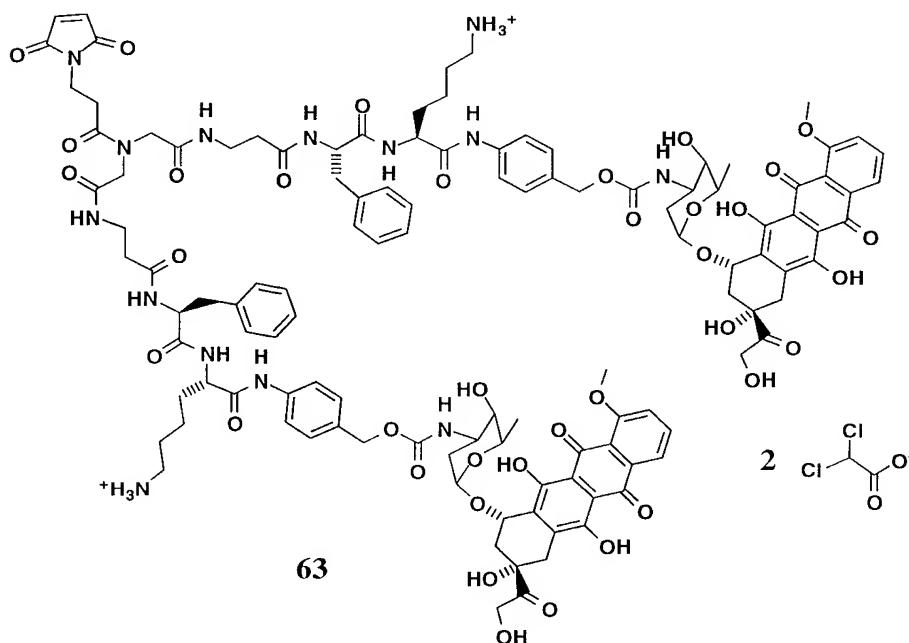
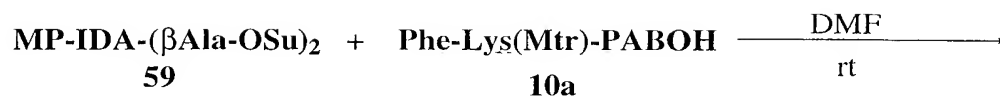
Scheme 8



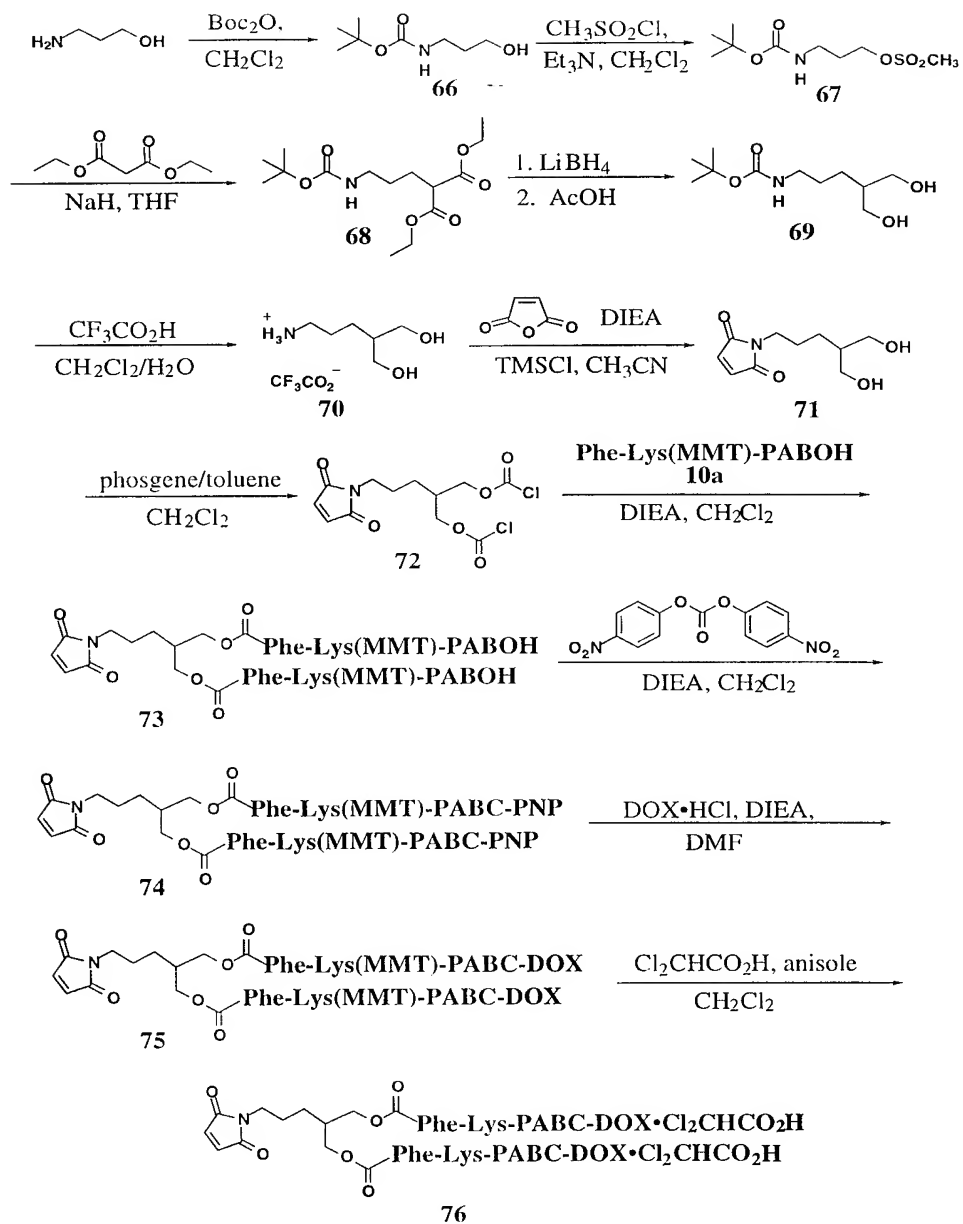
Scheme 9 (Part I)



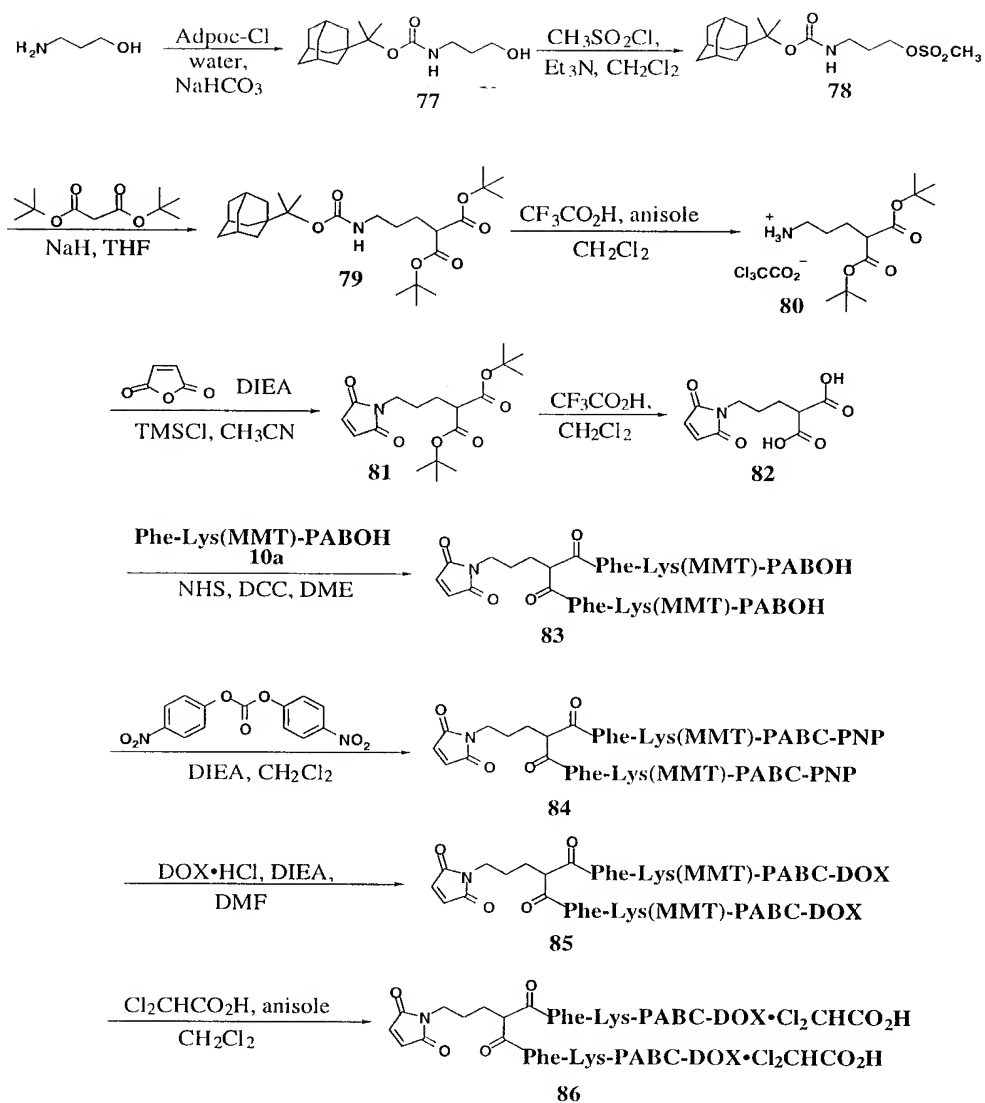
Scheme 9 (Part II)



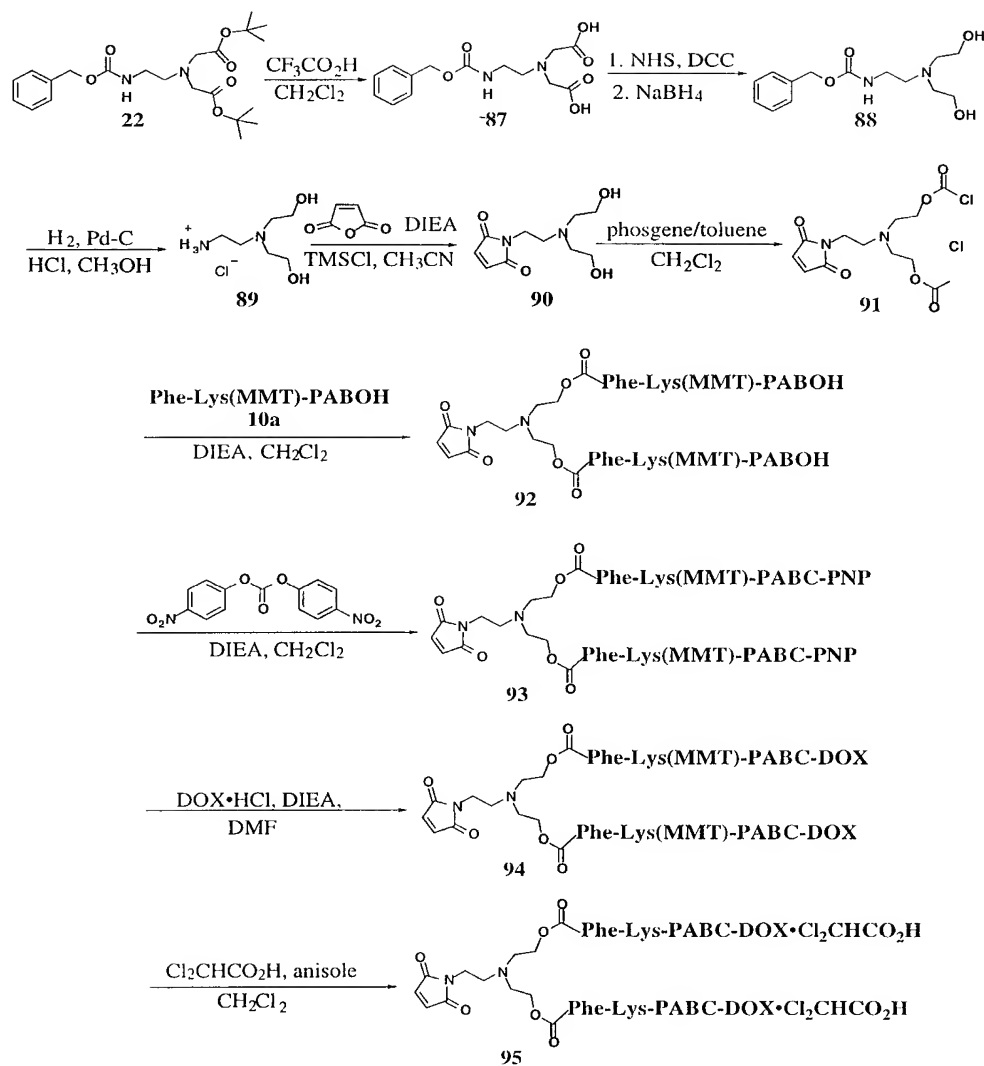
Scheme 10



Scheme 11



Scheme 12



The following examples are to illustrate the invention but should not be construed as a limitation thereon.

Example 1**Fmoc-Lys(Mtr)**

A stirred suspension of Fmoc-Lys hydrochloride (23.78 g, 56.42 mmol) in dry methylene chloride (250 mL) under argon at rt was treated with trimethylsilyl chloride (15 mL, 2.1 equiv) and DIEA (10.3 mL, 1.05 equiv). The mixture
5 was heated at reflux for 1 h, during which time it became homogeneous, and then cooled to 0°C. DIEA (31 mL, 3.1 equiv) was added, followed by p-anisylidiphenylmethyl chloride (18.29 g, 1.05 equiv). The reaction was stirred at rt for 14 h. The solvent was evaporated and the residue partitioned
10 between ethyl acetate and pH 5 buffer (0.05 M biphthalate). The organic phase was washed with more pH 5 buffer, water and brine, dried over sodium sulfate and evaporated to give a pale-yellow foam (34.71 g, 96%).
¹H-NMR (CDCl₃) δ 1.26 and 1.68 (m, 2H and 4H), 2.45 (m, 2H), 3.71 (s, 3H), 4.05-4.40 (m, 4H), 6.81 (d, 2H), 7.15-7.77 (m, 20H). MS (FAB) 641
15 (MH)⁺, 663 (M+Na)⁺, 679 (M+K)⁺. HRMS Calcd: 641.3015. Found: 641.3001.

Example 2**Lys(Mtr)**

Fmoc-Lys(Mtr) **1** (5.25 g, 8.19 mmol) in 1:1 methylene chloride/acetonitrile (80 mL) at rt was treated with diethylamine (80 mL). After 1.5 h the solvents were evaporated. The residue was flushed with acetonitrile (2 x 50 mL) at 60°C, and then triturated with ether (80 mL). The resulting solid was collected by filtration, washed with ether, and then dissolved as far as
25 possible in 1:1 methylene chloride/methanol. Some solid byproduct was removed by filtration and the filtrate was concentrated in vacuo. The resulting light tan solid was dried in vacuo for 4 h (3.2221 g, 94%). ¹H-NMR (DMSO-d₆) δ 1.34, 1.57 and 1.72 (m, 6H), 2.05 (m, 2H), 3.38 (m, 1H), 3.68 (s, 3H), 3.71 (d, 2H), 7.03-7.40 (m, 12H). MS (FAB) 419.2 (MH)⁺, 441.4 (M+Na)⁺,
30 457.4 (M+K)⁺. MS (FAB) 419.2 (MH)⁺, 441.4 (M+Na)⁺, 457.4 (M+K)⁺.

Example 3

Fmoc-Lys(Mtr)-OSu

A stirred solution of Fmoc-Lys(Mtr) **2** (5.5567 g, 8.672 mmol) and N-hydroxysuccinimide (1.0978 g, 1.1 equiv) in DME (100 mL) at 0°C was
5 treated with 0.5M DCC in methylene chloride (19.1 mL, 1.1 equiv). The mixture was gradually warmed to rt. after 14 h the solid DCU was filtered off and the filtrate evaporated to dryness. The crude active ester was used without further purification. ¹H-NMR (CDCl₃) δ 1.52, 1.73 and 1.92 (m, 6H),
2.14 (t, 2H), 2.79 (brs, 4H), 3.75 (s, 3H), 4.21 (t, 1H), 4.42 (d, 2H), 4.71 (m,
10 1H), 5.40 (d, 1H), 6.79 (d, 2H), 7.10-7.76 (m, 20H).

Example 4

Fmoc-Lys(Mtr)-Lys(Mtr)

A stirred solution of Fmoc-Lys(Mtr)-OSu **3** (ca 8.672 mmol) and Lys(Mtr) **2**
15 (3.6294 g, 1 equiv) in DMF (70 mL) at rt was treated with DIEA (4.5 mL, 3 equiv). After 4 h the mixture was diluted with ethyl acetate (250 mL). The solution was washed with pH 4 buffer (0.05M biphthalate) (2x), water and brine, dried over sodium sulfate and evaporated. the residue was flushed with methylene chloride (400 mL) to give a pale-yellow foam which was
20 carried on without further purification. ¹H-NMR (CDCl₃/CD₃OD) δ 1.10-1.85 (m, 12H), 2.22 (m, 4H), 3.70 (m, 6H), 4.21 (m, 5H), 6.76 (m, 4H), 7.00-7.75 (m, 32H). MS (ESI) 1041.6 (MH)⁺.

Example 5

Fmoc-Lys(Mtr)-Lys(Mtr)-PABOH

A stirred solution of Fmoc-Lys(Mtr)-Lys(Mtr) **4** (ca 8.672 mmol) and di-t-butylidicarbonate (2.8390 g, 1.5 equiv) in methylene chloride (100 mL) at rt was treated with pyridine (0.736 mL, 1.05 equiv). After 20 min p-aminobenzyl alcohol (1.6020 g, 1.5 equiv) was added. Stirring was
30 continued for 16 h and then the solvent was evaporated and the residue dried in vacuo for 2 h. The crude product was carried on without further purification. ¹H-NMR (CDCl₃/CD₃OD) δ 1.00-1.75 (m, 12H), 2.09 (m, 4H), 3.73 (m, 6H), 4.14 (m, 3H), 4.39 (m, 2H), 4.58 (m, 2H), 6.76 (m, 4H), 7.05-

7.75 (36H). MS (ESI) 1146.6 (MH)⁺. Anal. Calcd for C₇₄H₇₅N₅O₇-2H₂O: —
C-75.17, H-6.73, N-5.92. Found: C-74.89, H-6.37, N-5.75.

Example 6

5 **Lys(Mtr)-Lys(Mtr)-PABOH**

Fmoc-Lys(Mtr)-Lys(Mtr)-PABOH **5** (ca 8.672 mmol) in acetonitrile (100 mL) at
rt was treated with diethylamine (75 mL). After 2 h the solvents were
removed on the rotovap at 40°C and the residue dried in vacuo for 1 h and
then dissolved as far as possible in methylene chloride (150 mL). The solid
10 was removed by filtration and the filtrate concentrated in vacuo. The residue
was chromatographed on silica, eluting with 1) 2.5%, 2) 3%, and 3.5%
methanol/methylene chloride, to give the product as a colorless foam
(2.8403 g, 35% (4 steps)). ¹H-NMR (CDCl₃) δ 1.42 and 1.78 (m, 12H), 2.12
(brt, 4H), 3.32 (m, 1H), 3.75 (s, 3H), 3.79 (s, 3H), 4.44 (m, 1H), 4.57 (s, 2H),
15 6.77 (ABq, 4H), 7.10-7.70 (m, 28H), 7.91 (d, 1H), 9.09 (s, 1H). MS (ESI)
924.5 (MH)⁺. Anal. Calcd for C₅₉H₆₅N₅O₅-H₂O: C-75.21, H-7.17, N-7.43.
Found: C-75.37, H-6.96, N-7.24.

Example 7a

20 **Fmoc-Phe-OSu**

Fmoc-Phe (5.1043 g, 13.17 mmol) and NHS (1.592 g, 1.05 equiv) in
methylene chloride (100 mL) at 0°C were treated with DCC (2.854 g, 1.05
equiv). The ice bath was allowed to warm to rt and the mixture was stirred
for 14 h. DCU was removed by filtration and the filtrate was evaporated. The
25 resulting crude product, a colorless glass, was used without further
purification.

Example 7b

Fmoc-Val-OSu 7b

30 This was prepared from Fmoc-Val (7.02 g, 20.7 mmol) as described above
for **7a**.

Example 7c**Fmoc-Ala-OSu**

This was prepared from Fmoc-Ala (5.0414 g, 15.31 mmol) as described above for **7a**.

5

Example 8a**Fmoc-Phe-Lys(Mtr)**

A suspension of Lys(Mtr) **2** (4.686 g, 11.20 mmol) and NaHCO₃ (941.0 mg, 1 equiv) in water (100 mL) and DME (50 mL) was treated with a solution of Fmoc-Phe-OSu **7a** (11.20 mmol) in DME (50 mL). THF (25 mL) was then added to aid solubility. The mixture was stirred at rt for 2 days and then as much DME as possible was removed on the rotovap (bath at 30°C). The resulting gummy suspension was partitioned between ethyl acetate and pH 5 buffer. The organic phase was washed with water and brine, dried and evaporated to give a pale yellow foam. This was flushed with methylene chloride (100 mL). TLC showed the product to be satisfactorily pure and it was carried on without further purification (8.559 g, 97%). ¹H-NMR (CDCl₃/CD₃OD) δ 1.10-1.93 (m, 6H), 2.31 (t, 2H), 3.00 (m, 2H), 3.71 (s, 3H), 4.02-4.48 (m, 5H), 6.79 (d, 2H), 7.00-7.75 (m, 25H). MS (FAB) 788.2 (MH)⁺, 810.4 (M+Na)⁺, 826 (M+K)⁺. Anal. Calcd for C₅₀H₄₉N₃O₆·H₂O: C-74.51, H-6.38, N-5.21. Found: C-74.17, H-6.57, N-5.41.

15
20**Example 8b****Fmoc-Val-Lys(Mtr)**

This was prepared from Fmoc-Val-OSu **7b** (20.7 mmol) and Lys(Mtr) **2** (9.09 g, 1.05 equiv) as described above for **8a** (15.28 g, 100%). ¹H-NMR (CDCl₃/CD₃OD) δ 0.85 (m, 6H), 1.20 (m, 2H), 1.59 (m, 4H), 1.99 (m, 1H), 2.41 (m, 2H), 3.71 (s, 3H), 4.21 (m, 5H), 6.78 (d, 2H), 7.29 (m, 16H), 7.51 (brt, 2H), 7.71 (d, 2H). MS (FAB) 740 (MH)⁺, 762 (M+Na)⁺. HRMS Calcd: 740.3700. Found: 740.3712.

25
30

Example 8c**Fmoc-Ala-Lys(Mtr)**

A solution of Fmoc-Ala-OSu **7c** (15.31 mmol) in DME (100 mL) was added to a stirred solution of Lys(Mtr) **2** (6.4080 g, 1 equiv) and potassium bicarbonate (1.5331 mg, 1 equiv) in 2:1 water/DME (150 mL). After 16 h the DME was removed in vacuo and the resulting suspension was treated with citric acid (3.22 g, 1.1 equiv) in water (25mL). The mixture was extracted with ethyl acetate. The organic phase was washed with water (2x) and brine, dried over sodium sulfate, and concentrated in vacuo, giving a yellow foam which was carried on without further purification. ¹H-NMR (CDCl₃/CD₃OD) δ 1.25 (m, 2H), 1.32 (d, 3H), 1.57 (m, 3H), 1.73 (m, 1H), 2.41 (brt, 2H), 3.72 (s, 3H), 4.14 (m, 1H), 4.29 (m, 4H), 6.79 (d, 2H), 7.22 (m, 16H), 7.38 (d, 4H), 7.52 (m, 2H), 7.72 (d, 2H). MS (FAB) 712.2 (MH)⁺, 750.2 (M+K)⁺.

Example 9a**Fmoc-Phe-Lys(Mtr)-PAB-OH**

A stirred solution of Fmoc-Phe-Lys(Mtr) **8a** (7.728 g, 9.808 mmol) and p-aminobenzyl alcohol (1.450 g, 1.2 equiv) in methylene chloride (100 mL) at rt was treated with EEDQ (3.640 g, 1.5 equiv). After 20 h the solvent was evaporated (water bath at 30°C). The solid residue was triturated with ether (200 mL) and the resulting suspension sonicated for 15 min and left to stand at rt for 2 h. The resulting solid was collected by filtration, washed well with ether, and dried in vacuo (7.6140 g, 87%). ¹H-NMR (CDCl₃/CD₃OD) δ 0.98-1.91 (m, 6H), 2.06 (t, 2H), 2.97 (m, 2H), 3.71 (s, 3H), 4.12 (t, 1H), 4.20-4.41 (m, 4H), 4.59 (s, 2H), 6.72 (d, 2H), 7.00-7.73 (m, 29H). MS (FAB) 891.4 (MH)⁺, 916.7 (M+Na)⁺, 931 (M+K)⁺. Anal. Calcd for C₅₇H₅₆N₄O₆·H₂O: C-75.14, H-6.42, N-6.15. Found: C-75.25, H-6.02, N-6.49.

Example 9b**Fmoc-Val-Lys(Mtr)-PABOH**

This was prepared from Fmoc-Val-Lys(Mtr) **8b** (15.28 g, 20.65 mmol) as described above for **9a** (14.24 g, 82%). ¹H-NMR (DMSO-d₆) δ 0.84 (ABq, 6H), 1.15-1.80 (m, 6H), 1.94 (m, 3H), 2.37 (brt, 1H), 3.69 (s, 3H), 3.91 (t, 1H), 4.21 (m, 4H), 4.41 (d and m, 3H), 5.10 (t, 1H), 6.79 (d, 2H), 7.29 (m, 18H), 7.53 (d, 2H), 7.68 (t, 2H), 7.86 (m, 2H), 8.02 (d, 1H), 9.94 (brs, 1H). IR (KBr) 700, 740, 1650, 1695. MS (ESI) 845.6 (MH)⁺. Anal Calcd for C₅₃H₅₆N₄O₆·0.5H₂O: C-73.76, H-6.77, N-6.49. Found: C-74.01, H-6.68, N-6.63.

Example 9c**Fmoc-Ala-Lys(Mtr)-PABOH**

This was prepared from Fmoc-Ala-Lys(Mtr) **8c** (10.8985 g, 15.31 mmol) as described above for **9a** (11.0073 g, 88%). ¹H-NMR (CDCl₃/CD₃OD) δ 1.29 (d and m, 5H), 1.42 (m, 2H), 1.57 (m, 1H), 1.79 (m, 1H), 2.05 (m, 2H), 3.68 (s, 3H), 4.11 (m, 2H), 4.30 (m, 3H), 4.51 (s, 2H), 6.69 (d, 2H), 7.00-7.55 (m, 22H), 7.66 (d, 2H). MS (ESI) 818.0 (MH)⁺. Anal Calcd for C₅₁H₅₂N₄O₆·H₂O: C-73.36, H-6.52, N-6.71. Found: C-73.02, H-6.72, N-6.54.

Example 10a**Phe-Lys(Mtr)-PAB-OH**

Fmoc-Phe-Lys(Mtr)-PAB-OH **9a** (4.2857 g, 4.80 mmol) in methylene chloride (35 mL) at rt was treated with diethylamine (50 mL). The mixture was sonicated briefly and stirred at rt for 4 h. after which time no starting material was observed by TLC. The solvents were evaporated and the residue was flushed with methylene chloride and chromatographed on silica, eluting with 1) 2%, 2) 3%, and 3) 4% methanol/methylene chloride, to give the product as a colorless foam (2.230 g, 69%). ¹H-NMR (CDCl₃) δ 1.26-2.00 (m, 6H), 2.12 (t, 2H), 2.75 and 3.21 (ABq, each 1H), 3.68 (ABq, 1H), 3.76 (s, 3H), 4.42 (q, 1H), 4.66 (brs, 2H), 6.79 (d, 2H), 7.10-7.42 (m, 21H), 7.81 (d, 1H), 8.71 (s,

1H). MS (FAB) 693.4 (M+Na)⁺, 709 (M+K)⁺. Anal. Calcd for C₄₂H₄₆N₄O₄·
1/2H₂O: C-74.20, H-6.97, N-8.24. Found: C-74.28, H-7.00, N-8.34.

Example 10b

5 **Val-Lys(Mtr)-PABOH**

A solution of Fmoc-Val-Lys(Mtr)-PABOH **9b** (13.94 g, 16.50 mmol) in 5:1 methylene chloride/methanol (75 mL) was treated with diethylamine (150 mL). The reaction was sonicated for 30 min and then allowed to stand at rt for 3 h. The solvents were evaporated and the residue flushed with
10 methylene chloride. The remaining material was chromatographed on silica, eluting with 1) 2%, 2) 3%, 3) 3.5%, and 4) 4% methanol/methylene chloride, to give the product as a colorless foam (6.2510 g, 61%). ¹H-NMR (CDCl₃/CD₃OD) δ 0.80 (d, 3H), 0.97 (d, 3H), 1.48 (m, 4H), 1.70 (m, 1H), 1.90 (m, 1H), 2.11 (t, 2H), 2.27 (m, 1H), 3.27 (d, 1H), 3.76 (s, 3H), 4.53 (q,
15 1H), 4.58 (s, 2H), 6.78 (d, 2H), 7.10-7.50 (m, 16H), 7.95 (d, 1H), 9.11 (s, 1H). IR (KBr) 706, 830, 1034, 1248, 1510, 1648, 3310. MS (ESI) 623.2 (MH)⁺. HRMS Calcd for C₃₈H₄₇N₄O₄: 623.3598. Found: 623.3606. Anal. Calcd for C₃₈H₄₆N₄O₄·H₂O: C, 71.22, H, 7.55, N, 8.74. Found: C, 71.41, H, 7.51, N, 8.66.

20

Example 10c

Ala-Lys(Mtr)-PABOH

A solution of Fmoc-Ala-Lys(Mtr)-PABOH **9c** (11.0073 g, 13.47 mmol) in DMF (50 mL) at rt was treated with diethylamine (25 mL). After 2.5 h the mixture
25 was concentrated under high vacuum and the residue was flushed with methylene chloride (2x100 mL) and then toluene (150 mL). The remaining material was chromatographed on silica, eluting with 1) 4:1, 2) 1:1, 3) 1:1.6 and 4) 1:3 methylene chloride/(5:1 ethyl acetate/methanol), to give the product as a pale-yellow foam (5.7817 g, 72%). ¹H-NMR (CDCl₃/CD₃OD) δ
30 1.32 (d, 3H), 1.25-2.01 (m, 6H), 2.13 (t, 2H), 3.51 (q, 1H), 3.75 (s, 3H), 4.48 (q, 1H), 4.60 (brs, 2H), 6.79 (d, 2H), 7.12-7.50 (m, 16H), 7.89 (d, 1H), 8.89 (brs, 1H). MS (ESI) 595.2 (MH)⁺.

Example 11**N, N-Bis-Carbobenzyloxyethylenediamine**

Benzyl chloroformate (40 mL, 280 mmol) was added dropwise over 45 min to a vigorously stirred solution of ethylenediamine (9.4 mL, 0.5 equiv) in 1M NaOH (310 mL, 1 equiv) at 0°C. A heavy white precipitate formed and the mixture was stirred for 4 h following addition. The resulting solid product was collected by filtration, washed with water and then hexane, and then air-dried overnight (46.17 g, 100%). ¹H-NMR (CDCl₃) δ 3.32 (s, 4H), 5.10 (s, 4H), 5.23 (br, 2H), 7.35 (s, 10H). MS (ESI) 329.3 (MH)⁺.

Example 12**N-Carbobenzyloxyethylenediamine hydrochloride**

N, N-Bis-Carbobenzyloxyethylenediamine **11** (23.91 g, 72.8 mmol) in glacial acetic acid (100 mL) was treated with 12M HCl (12.1 mL, 2 equiv). The stirred mixture was heated at reflux for 1 h and then left to stand at rt overnight. A small amount of solid was removed by filtration and the filtrate was diluted with ether (700 mL) and left to stand at rt for 2 h. The resulting white solid product was collected by filtration, washed repeatedly with ether, and dried in a vacuum dessicator overnight (9.2440 g, 55%). ¹H-NMR (DMSO-d₆) δ 2.87 (t, 2H), 3.30 (q, 2H), 5.04 (s, 2H), 7.33 (m, 5H), 7.49 (brt, 1H), 8.22 (br, 3H).

Example 13**N-Carbobenzyloxy-N', N'-bis-(t-butoxycarbonyl)ethylenediamine**

A stirred solution of N-carbobenzyloxyethylenediamine hydrochloride **12** (5.0009 g, 21.68 mmol) in 3:1 methanol/methylene chloride (100 mL) at rt was treated with t-butyl acrylate (64 mL, 20 equiv) and triethylamine (3.3 mL, 1.1 equiv). After 2 days the solvents were removed on the rotovap. The residue was dried in vacuo for 30 min and then partitioned between 30% ethyl acetate/ether and 50% sat. NaHCO₃. The organic phase was washed with water and brine, dried over sodium sulfate and evaporated. The

residue was chromatographed on silica, eluting with 20% ethyl acetate/hexane, to give the product as a thick, colorless oil (9.3801 g, 96%).

$^1\text{H-NMR}$ (CDCl_3) δ 1.42 (s, 18H), 2.33 (t, 4H), 2.54 (t, 2H), 2.72 (t, 4H), 3.28 (q, 2H), 5.11 (s, 2H), 5.49 (brt, 1H), 7.33 (m, 5H). MS(DCI) 451 (MH) $^+$. Anal.

5 Calcd for $\text{C}_{24}\text{H}_{38}\text{N}_2\text{O}_6 \cdot 0.5 \text{H}_2\text{O}$: C, 62.72, H, 8.55, N, 6.09. Found: C, 62.54, H, 8.27, N, 5.95.

Example 14

N, N-Bis-(t-butoxycarbonylethyl)ethylenediamine dihydrochloride

10

A solution of N-carbobenzyloxy-N', N'-bis-t-butoxypropionylethylenediamine **13** (18.3640 g, 40.76 mmol) was passed through a bed of Raney nickel in a fritted glass funnel. The filtrate was degassed with nitrogen and 10% Pd-C (1 g) and 12M HCl (6.8 mL, 2 equiv) were added. The mixture was

15 hydrogenated on a Parr shaker at 50 psi for 16 h and then degassed once again with nitrogen. The catalyst was filtered off on a bed of celite. The filtrate was concentrated in vacuo and the residue flushed with methylene chloride (2x), giving an off-white foam (15.8696 g, 100%). $^1\text{H-NMR}$ (CDCl_3) δ 1.48 (s, 18H), 2.97 (brt, 4H), 3.53 (m, 4H), 3.79 (m, 2H), 3.88 (m, 2H), 8.61
20 (br, 3H), 11.24 (br, 1H).

Example 15

N-Maleoyl-N', N'-bis-(t-butoxycarbonylethyl)-ethylenediamine

25 A stirred suspension of N, N-bis-t-butoxypropionylethylenediamine dihydrochloride **14** (5.3363 g, 13.70 mmol) in acetonitrile (30 mL) at 0°C under argon was treated with DIEA (4.77 mL, 2 equiv) and maleic anhydride (1.4111 g, 1.05 equiv). The reaction was allowed to warm gradually to rt. After 16 h trimethylsilyl chloride (5.2 mL, 3 equiv) and DIEA (7.2 mL, 3 equiv)
30 were added. The mixture was heated at reflux for 4 h. Upon cooling, the solvents were removed in vacuo and the residue taken up in ethyl acetate. The solution was washed with sat. NaHCO_3 , water and brine, dried over sodium sulfate and evaporated. The residue was chromatographed on

silica, eluting with 2.5:1 hexane/ethyl acetate, to give the product as a pale-
yellow oil (2.9986 g, 55%). $^1\text{H-NMR}$ (CDCl_3) δ 1.42 (s, 18H), 2.30 (t, 4H),
2.61 (t, 2H), 2.73 (t, 4H), 3.59 (t, 2H), 6.68 (s, 2H). MS (ESI) 397.4 (MH) $^+$.
Anal. Calcd for $\text{C}_{20}\text{H}_{32}\text{N}_2\text{O}_6$: C-60.59, H-8.13, N-7.07. Found: C-60.65, H-
5 7.98, N-6.88.

Example 16

N-Maleoyl-N', N'-bis-(carboxyethyl)ethylenediamine p-toluenesulfonic acid

10 N-Maleoyl-N', N'-bis-t-butoxypropionylethylenediamine **15** (3.9934 g, 10.07
mmol) in methylene chloride (100 mL) at rt was treated with p-
toluenesulfonic acid monohydrate (5.75 g, 3 equiv). The reaction was stirred
at rt overnight. The solvent was evaporated at rt and the gummy residue
dried in vacuo for 3 h. NMR showed complete loss of t-butyl ester signal.
15 The crude product was carried on without further purification. $^1\text{H-NMR}$
(DMSO- d_6) δ (contains 3 equiv of p-toluenesulfonic acid) 2.29 (s, 9H), 2.76
(m, 4H), 3.39 (m, 6H), 3.81 (brt, 2H), 7.11 (s, 2H), 7.17 (d, 6H), 7.49 (d, 6H),
9.27 (br, 1H).

Example 17

N-Maleoyl-N', N'-bis-(succinimidylloxycarbonyl- ethyl)ethylenediamine

A stirred solution of N-maleoyl-N', N'-bis-propionylethylenediamine p-
toluenesulfonate **16** (ca 10.07 mmol, containing two extra equiv of p-
25 toluenesulfonic acid), triethylamine (4.4 mL, 3.1 equiv) and N-
hydroxysuccinimide (2.5502 g, 2.2 equiv) in acetonitrile (90 mL) at 0°C was
treated with 0.5M DCC in methylene chloride (45.3 mL, 2.25 equiv). After 14
h the reaction was concentrated in vacuo and the residue partitioned
between ethyl acetate and water. The aqueous phase was re-extracted with
30 ethyl acetate. The combined organic phases were washed with water (2x)
and brine, dried over sodium sulfate and evaporated. The residue was
chromatographed on silica, eluting with 1:1 ethyl acetate/methylene chloride,

to give the product as a pale-yellow foam (1.2043 g, 25%, 2 steps). $^1\text{H-NMR}$ (CDCl₃) δ 2.72 (t, 4H), 2.84 (brs, 6H), 2.92 (t, 4H), 3.61 (t, 2H), 6.68 (s, 2H).

Example 18a

5 **MEt-IDP-(Lys(Mtr)-Lys(Mtr)-PABOH)₂**

A solution of Lys(Mtr)-Lys(Mtr)-PABOH **6** (2.0903 g, 2.262 mmol) and N-maleoyl-N', N'-bis-succinimidylxypropionylethylenediamine **17** (541.0 mg, 0.5 equiv) in DME (30 mL) was stirred for 2 days at rt. The reaction was concentrated in vacuo at 30°C and the residue dissolved in ethyl acetate.

- 10 The solution was washed with water (2x) and brine, dried over sodium sulfate and evaporated to give the product as a colorless glass (2.3623 g, 99%). $^1\text{H-NMR}$ (CDCl₃/CH₃OD) δ 1.10-1.97 (m, 24H), 2.09 (m, 8H), 2.48 (m, 2H), 2.63 (m, 4H), 3.45 (m, 2H), 3.72 (s, 6H), 3.74 (s, 6H), 4.31 (m, 2H), 4.46 (m, 2H), 4.50 (s, 4H), 6.52 (s, 2H), 6.77 (m, 8H), 7.05-7.52 (m, 56H). MS
- 15 (ESI) 2096.4 (MH)⁺.

Exampel 18b**MEt-IDP-(Phe-Lys(Mtr)-PABOH)₂**

A stirred mixture of N-maleoyl-N', N'-bis-(carboxyethyl)ethylenediamine p-toluenesulfonic acid **16** (1.0643 g, 1.696 mmol) in DME (50 mL) at 0°C was
5 treated with triethylamine (0.6 mL, 2.5 equiv), Phe-Lys(Mtr)-PABOH **10a** (2.2750 g, 2 equiv), NHS (439.0 mg, 2.25 equiv) and DCC (0.5M in methylene chloride, 8.5 mL, 2.5 equiv). The reaction was allowed to warm gradually to rt. After 16 h the DCU byproduct was filtered off and the filtrate was concentrated in vacuo. The residue was triturated with ether and the
10 resulting solid was collected by filtration, washed with ether and dried in vacuo (2.0494 g, 76%). ¹H-NMR (CDCl₃/CD₃OD) δ 1.20-1.95 (m, 12H), 2.07 (brt, 4H), 2.18 (m, 4H), 2.39 (t, 2H), 2.54 (m, 4H), 3.02 (m, 4H), 3.29 (m, 2H), 3.71 (s, 6H), 4.41 (m, 2H), 4.56 (brs, 4H), 4.69 (m, 2H), 6.57 (s, 2H), 6.72 (d, 4H), 6.97-7.55 (m, 42H), 9.00 (br, 2H). MS(FAB) 1611 (M+Na)⁺, 1627
15 (M+K)⁺.

Example 18c**MEt-IDP-(Ala-Lys(Mtr)-PABOH)₂**

This was prepared from N-maleoyl-N', N'-bis-succinimidylxy-propionylethylenediamine **17** (. ¹H-NMR (CDCl₃/CD₃OD) δ 1.20-1.80 (m,
20 15H), 1.95 (m, 4H), 2.13 (m, 4H), 2.39 (m, 2H), 2.55 (m, 4H), 3.31 (m, 2H), 3.60 (s, 6H), 4.28 (m, 4H), 4.40 (brs, 4H), 6.50 (s, 2H), 6.61 (d, 4H), 6.90-7.50 (m, 32H). MS (ESI) 1437.8 (MH)⁺. HRMS Calcd for C₈₄H₉₆N₁₀O₁₂Na: 1459.7107. Found: 1459.707.

25

Example 19a**MEt-IDP-(Lys(Mtr)-Lys(Mtr)-PABC-PNP)₂**

MEt-IDP-(Lys(Mtr)-Lys(Mtr)-PABOH)₂ **18a** (2.3623 g, 1.127 mmol) and bis-p-nitrophenylcarbonate (1.5424 g, 4.5 equiv) in dry methylene chloride (60
30 mL) were treated with freshly activated 4Å powdered sieves (8.5 g) and DIEA (0.883 mL, 4.5 equiv). The mixture was stirred at rt for 16 h and then filtered. The filtrate was concentrated in vacuo to give a yellow solid which was

dissolved as far as possible in ether (200 mL). The suspension was sonicated and stirred overnight at rt. The resulting white solid was collected by filtration, washed repeatedly with ether, and dried in vacuo (2.4084 g, 88%). ¹H-NMR (CDCl₃/CH₃OD) δ 1.10-1.97 (m, 24H), 2.04 (m, 8H), 2.28 (m, 4H), 2.48 (m, 2H), 2.62 (m, 4H), 3.41 (m, 2H), 3.71 (s, 6H), 3.73 (s, 6H), 4.36 (m, 2H), 4.42 (m, 2H), 5.14 (s, 4H), 6.51 (s, 2H), 6.73 (m, 8H), 7.00-7.60 (m, 60H), 8.20 (d, 4H). Anal. Calcd for C₁₄₄H₁₄₈N₁₄O₂₂-2H₂O: C-70.23, H-6.22, N-7.96. Found: C-70.11, H-6.22, N-7.96.

10

Example 19b**MEt-IDP-(Phe-Lys(Mtr)-PABC-PNP)₂**

This was prepared from MEt-IDP-(Phe-Lys(Mtr)-PABOH)₂ **18b** (517.8 mg, 0.326 mmol) as described above for **19a** (509.1 mg, 74%). ¹H-NMR (CDCl₃/CD₃OD) δ 1.20-1.97 (m, 12H), 2.08 (m, 4H), 2.24 (m, 4H), 2.40 (t, 2H), 2.61 (m, 4H), 3.10 (m, 4H), 3.29 (m, 2H), 3.73 (s, 6H), 4.41 (q, 2H), 4.69 (q, 2H), 5.22 (s, 4H), 6.58 (s, 2H), 6.77 (d, 4H), 7.00-7.50 (m, 42H), 7.56 (d, 4H), 8.22 (d, 4H), 8.74 (brs, 2H).

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Example 19c**MEt-IDP-(Ala-Lys(Mtr)-PABC-PNP)₂**

This was prepared from MEt-IDP-(Ala-Lys(Mtr)-PABOH)₂ **18c** (6.24 g, 4.34 mmol) as described above for **19a** (6.00 g, 74%). ¹H-NMR (CDCl₃/CD₃OD) δ 1.26 (d, 6H), 1.33 (m, 4H), 1.42 (m, 4H), 1.62 (m, 2H), 1.83 (m, 2H), 2.18 (m, 4H), 2.41 (m, 6H), 2.64 (m, 4H), 3.72 (s, 6H), 4.40 (m, 4H), 5.18 (brs, 4H), 6.60 (s, 2H), 6.72 (d, 4H), 7.05-7.50 (m, 38H), 7.56 (d, 4H), 8.21 (d, 4H).

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Example 20a**MEt-IDP-(Lys(Mtr)-Lys(Mtr)-PABC-DOX)₂**

A stirred solution of MEt-IDP-(Lys(Mtr)-Lys(Mtr)-PABC-PNP)₂ **19a** (753.0 mg, 0.3103 mmol) and DOX hydrochloride (377.9 mg, 2.1 equiv) in DMF (25 mL) at rt was treated with DIEA (0.114 mL, 2.1 equiv). After 3 days the mixture was poured into ethyl acetate (200 mL) and the solution was washed

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with water (4x) and brine, dried over sodium sulfate, and evaporated to give an orange glass. This was triturated with ether (200 mL), and the resulting solid was collected by filtration and then chromatographed on silica, eluting with 16:1 methylene chloride/methanol, to give the product as an orange solid (803.8 mg, 80%). ¹H-NMR (DMF-d₇) δ 1.23 (d, 6H), 1.30-1.90 (m, 28H), 2.04 (m, 10H), 2.30 (m, 6H), 2.41 (d, 2H), 2.51 (m, 2H), 2.72 (m, 4H), 3.12 (q, 4H), 3.64 (m, 2H), 3.75 (s, 12H), 3.91 (m, 2H), 4.08 (s and m, 8H), 4.33 (m, 4H), 4.45 (m, 2H), 4.79 (s, 4H), 4.89 (d, 2H), 4.96 (brs, 4H), 5.12 (m, 2H), 5.40 (brs, 2H), 5.60 (s, 2H), 6.72 (d, 2H), 6.87 (q, 8H), 6.92 (s, 2H), 7.10-7.50 (m, 54H), 7.67 (m, 6H), 7.93 (m, 2H), 9.91 (brs, 2H).

Example 20b

MEt-IDP-(Phe-Lys(Mtr)-PABC-DOX)₂

This was prepared from MEt-IDP-(Phe-Lys(Mtr)-PABC-PNP)₂ **19b** (340.8 mg, 0.161 mmol) as described above for **20a** (411.2 mg, 93%). ¹H-NMR (CDCl₃/CD₃OD) δ 1.10-1.65 (m and d, 16H), 2.07 (m, 6H), 2.10-2.45 (m, 10H), 2.55 (m, 4H), 3.00 (m, 6H), 3.26 (m and d, 4H), 3.71 (s, 6H), 3.82 (m, 2H), 4.00 (brs, 6H), 4.11 (m, 2H), 4.40 (m, 2H), 4.63 (m, 2H), 4.74 (s, 4H), 4.90 (ABq, 4H), 5.13 (brs, 2H), 5.44 (brs, 2H), 5.58 (brd, 2H), 6.60 (s, 2H), 6.71 (d, 4H), 6.90-7.55 (m, 44H), 7.72 (t, 2H), 7.94 (d, 2H).

Example 20c

MEt-IDP-(Ala-Lys(Mtr)-PABC-DOX)₂

This was prepared from MEt-IDP-(Ala-Lys(Mtr)-PABC-PNP)₂ **19c** (6.00 g, 3.39 mmol) as described above for **20a** (620 mg, 7%). ¹H-NMR (CDCl₃/CD₃OD) δ 1.17 (m, 16H), 1.29 (m, 4H), 1.40 (m, 4H), 1.72 (m, 7H), 1.99 (m, 4H), 2.15 (m, 10H), 2.53 (m, 6H), 3.03 (ABq, 4H), 3.39 (m, 2H), 3.52 (brs, 2H), 3.64 (s, 6H), 3.95 (s, 6H), 4.04 (m, 2H), 4.26 (m, 2H), 4.33 (m, 2H), 4.68 (s, 4H), 4.80 (ABq, 4H), 5.10 (brs, 2H), 5.47 (brs, 2H), 6.55 (s, 2H), 6.64 (d, 4H), 6.98-7.40 (m, 34H), 7.68 (t, 2H), 7.89 (d, 2H).

Example 21a**MEt-IDP-(Lys-Lys-PABC-DOX)₂•5Cl₂CHCO₂H**

A stirred solution of MEt-IDP-(Lys(Mtr)-Lys(Mtr)-PABC-DOX)₂ **20a** (318.7 mg, 0.0985 mmol) in methylene chloride (21 mL) at rt was treated with
5 anisole (3.2 mL, 300 equiv) and dichloroacetic acid (0.244 mL, 30 equiv).

After 2.5 h ethyl acetate (50 mL) was added. The mixture was stored at 4°C for 2 h. The resulting solid was collected by filtration and washed repeatedly with ethyl acetate followed by ether, and then air-dried (274.5 mg, 99.8 %).

¹H-NMR (DMF-d₇) δ 1.23 (d, 6H), 1.54 (m, 8H), 1.75 (m, 18H), 2.01 (m, 2H),
10 2.22 (q, 2H), 2.40 (m, 6H), 3.07 (m, 8H), 3.89 (m, 2H), 4.06 (s, 6H), 4.34 (m, 4H), 4.52 (m, 2H), 4.79 (s, 4H), 4.96 (s, 4H), 5.13 (brs, 2H), 5.39 (s, 2H), 5.72 (m, 2H), 6.30 (s, 5H), 6.77 (d, 2H), 6.99 (s, 2H), 7.30 (d, 4H), 7.71 (m, 6H), 7.93 (m, 2H), 8.45 (m, 8H). MS (ESI) 2147.6 (MH)⁺.

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Example 21b**MEt-IDP-(Phe-Lys-PABC-DOX)₂•3Cl₂CHCO₂H**

This was prepared from MEt-IDP-(Phe-Lys(Mtr)-PABC-DOX)₂ **20b** (280.0 mg, 0.103 mmol) as described above for **21a** (237.1 mg, 90%). ¹H-NMR

(CDCl₃/CD₃OD) δ 1.20 (d, 6H), 1.37 (m, 4H), 1.70 (m, 10H), 2.06 (m, 2H),
20 2.29 (m, 6H), 2.61 (m, 2H), 2.82 (m, 8H), 3.00 (m, 4H), 3.16 (d, 2H), 3.75 (m, 2H), 3.99 (s, 6H), 4.07 (m, 2H), 4.44 (m, 4H), 4.70 (s, 4H), 4.89 (ABq, 4H), 5.16 (brs, 2H), 5.40 (brs, 2H), 5.85 (s, 3H), 6.63 (s, 2H), 7.08 (m, 14H), 7.39 (d, 2H), 7.69 (t, 2H), 7.90 (d, 2H). MS (ESI) 1092.8 (MH₂)²⁺, 2184.8 (MH)⁺.

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Example 21c**MEt-IDP-(Ala-Lys-PABC-DOX)₂•3Cl₂CHCO₂H**

This was prepared from MEt-IDP-(Ala-Lys(Mtr)-PABC-DOX)₂ **20c** (569 mg, 0.248 mmol) as described above for **21a** (455 mg, 80%). ¹H-NMR

(CDCl₃/CD₃OD) δ 1.17 (d and m, 8H), 1.22 (brd, 6H), 1.38 (m, 4H), 1.65 (m,
30 10H), 2.02 (d, 2H), 2.23 (d, 2H), 2.41 (m, 4H), 2.82 (m, 6H), 2.99 (m, 4H), 3.10 (d, 2H), 3.54 (m and brs, 4H), 3.93 (s, 6H), 4.05 (m, 2H), 4.19 (m, 2H), 4.40 (m, 2H), 4.68 (brs, 4H), 4.82 (ABq, 4H), 5.10 (brs, 2H), 5.38 (brs, 2H),

5.79 (s, 3H), 6.61 (s, 2H), 7.09 (d, 4H), 7.29 (m, 2H), 7.40 (d, 4H), 7.67 (t, 2H), 7.82 (d, 2H). MS (ESI) 2032.8 (MH)⁺. MS (ESI) 1016.6 (MH₂)²⁺, 2032.4 (MH)⁺.

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Example 22**N-Carbobenzyloxy-N', N'-bis-(t-butoxycarbonylmethyl)-ethylenediamine**

A stirred mixture of N-carbobenzyloxyethylenediamine hydrochloride **12** (13.95 g, 60.48 mmol) and potassium bicarbonate (24.22 g, 4 equiv) in DMF (100 mL) at 0°C was treated with t-butyl bromoacetate (22 mL, 2.25 equiv), dropwise over 10 min. The mixture was allowed to warm to rt. Stirring was continued for 12 h and then the mixture was partitioned between 3:1 ether/ethyl acetate and 50% saturated sodium bicarbonate. The organic phase was washed with water (3x), brine, dried over sodium sulfate and evaporated to give a thick oil. This was flushed with heptane several times at 60°C to remove excess t-butyl bromoacetate and then chromatographed on silica, eluting with 1) hexane, 2) 5% ethyl acetate/hexane, and 3) 25% ethyl acetate/hexane, to give the product as a thick, colorless oil (18.53 g, 73%). ¹H-NMR (CDCl₃) δ 1.43 (s, 9H), 2.83 (t, 2H), 3.22 (q, 2H), 3.41 (s, 4H), 5.12 (s, 2H), 6.04 (brt, 1H), 7.33 (m, 5H). MS (ESI) 423.4 (MH)⁺.

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Example 23**N, N-Bis-(t-butoxycarbonylmethyl)ethylenediamine dihydrochloride**

A solution of N-carbobenzyloxy-N', N'-bis-(t-butoxyacetyl)ethylenediamine **22** (15.02 g, 35.54 mmol) in ethanol (75 mL) was filtered through a bed a Raney nickel. Acetic acid (6.1 mL, 3 equiv) was added and the mixture was hydrogenated as described above for **4**. the crude product was carried on without further purification.

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Example 24**N-Maleoyl-N', N'-bis-(t-butoxycarbonylmethyl)-
ethylenediamine**

N, N-Bis-(t-butoxyacetyl)ethylenediamine dihydrochloride **23** (ca. 35.54
5 mmol) in CHCl_3 (150 mL) at rt was treated with maleic anhydride (3.83 g, 1.1
equiv). The mixture was stirred for 14 h and then the solvent was
evaporated. The residue was flushed with CHCl_3 (200 mL) and then
dissolved in acetonitrile (200 mL) under argon. The stirred mixture was
cooled to 0°C and treated with trimethylsilyl chloride (18 mL, 4 equiv) and
10 triethylamine (20 mL, 4 equiv). The ice bath was removed and the mixture
was heated at reflux for 2.5 h. The solvents were removed under reduced
pressure and the residue partitioned between ether and 30% sat sodium
bicarbonate. The organic phase was washed with water, brine, dried and
evaporated. The residue was chromatographed on silica, eluting with 2.5:1
15 hexane/ethyl acetate, to give the product as a waxy solid (8.25 g, 63%, 3
steps). $^1\text{H-NMR}$ (CDCl_3) δ 1.42 (s, 9H), 2.91 (t, 2H), 3.41 (s, 4H), 3.62 (t,
2H), 6.68 (s, 2H). MS (DCI) 369 (MH^+).

Example 25**N-Maleoyl-N', N'-bis-(carboxymethyl)ethylenediamine•
p-toluenesulfonic acid**

A stirred solution of N-maleoyl-N', N'-bis-(t-butoxycarbonylmethyl)-
ethylenediamine **24** (2.6718 g, 7.25 mmol) in methylene chloride (70 mL) at
rt was treated with p-toluenesulfonic acid monohydrate (4.1400 g, 3 equiv).
25 After 36 h the resulting white solid was collected by filtration, washed
repeatedly with methylene chloride and ether, and then dried in vacuo. NMR
showed 1.4-1.5 equiv of p-toluenesulfonic acid per equiv of product (3.7531
g, 100%). $^1\text{H-NMR}$ (DMSO-d_6) δ 2.29 (s, 3.9H), 3.39 (t, 2H), 3.77 (t, 2H),
4.07 (s, 4H), 7.08 (s, 2H), 7.12 (d, 2.6H), 7.49 (d, 2.6H). MS (DCI) 257
30 (MH^+). Anal. Calcd for $\text{C}_{10}\text{H}_{13}\text{N}_2\text{O}_6 \cdot 1.3\text{C}_7\text{H}_7\text{O}_3\text{S} \cdot 2\text{H}_2\text{O}$: C-44.48, H-5.10,
N-5.43. Found: C-44.51, H-4.96, N-5.20.

Example 26a**MEt-IDA-(Phe-Lys(Mtr)-PABOH)₂**

A stirred solution of Phe-Lys(Mtr)-PABOH **10a** (2.5179 g, 3.753 mmol), N-maleoyl-N', N'-bis-(carboxymethyl)ethylenediamine • p-toluenesulfonic acid **25** (804.0 mg, 0.5 equiv) and N-hydroxysuccinimide (475.2 mg, 1 equiv) in DME (150 mL) at 0°C was treated with DIEA (0.33 mL, 0.5 equiv) and 0.5M DCC in methylene chloride (9.0 mL, 1.2 equiv). After 6 h DCU was removed by filtration and the filtrate was concentrated in vacuo. The residue was partitioned between ethyl acetate and water. The organic phase was washed with water (2x) and brine, dried over sodium sulfate, and concentrated in vacuo. The residue was dissolved as far as possible in methylene chloride (100 mL) and then the mixture was diluted with ether (100 mL). After being left to stand overnight the resulting solid was collected by filtration and air-dried (2.6295 g, 90%). ¹H-NMR (CDCl₃) δ 1.05-1.98 (m, 12H), 2.09 (brt, 4H), 2.51 (brt, 2H), 2.91 (brs, 4H), 3.03 (m, 4H), 3.39 (brt, 2H), 3.72 (s, 6H), 4.34 (q, 2H), 4.52 (s, 4H), 4.71 (q, 2H), 6.55 (s, 2H), 6.78 (d, 4H), 7.03-7.50 (m, 34H), 8.72 (brs, 2H). MS (FAB) 1583.9 (M+Na)⁺, 1600.9 (M+K)⁺.

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Example 26b**MEt-IDA-(Val-Lys(Mtr)-PABOH)₂**

This was prepared from N-maleoyl-N', N'-bis-(carboxymethyl)ethylenediamine•p-toluenesulfonic acid **25** (500 mg, 1.16 mmol) and Val-Lys(Mtr)-PABOH **10b** (1.44 g, 2 equiv) as described above for **26a** (1.68 g, 95%). ¹H-NMR (CDCl₃/CD₃OD) δ 0.90 (t, 12H), 1.10-1.88 (m, 14H), 2.10 (m, 4H), 2.71 (brt, 2H), 3.24 (brs, 4H), 3.46 (m, 2H), 3.77 (s, 6H), 4.25 (d, 2H), 4.42 (t, 2H), 4.58 (s, 4H), 6.63 (s, 2H), 6.79 (d, 4H), 7.05-7.55 (m, 32H). IR (KBr) 698, 1510, 1638, 1708. MS (ESI) 1466.0 (MH)⁺. Anal. Calcd for C₈₆H₁₀₀N₁₀O₁₂•2H₂O: C-68.78, H-6.98 N-9.33. Found: C-68.87, H-7.54, N-9.48.

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Example 27a**MEt-IDA-(Phe-Lys(Mtr)-PABC-PNP)₂**

A stirred suspension of MEt-IDA-(Phe-Lys(Mtr)-PABOH)₂ **26a** (471.5 mg, 0.319 mmol) and bis-p-nitrophenylcarbonate (971.0 mg, 10 equiv) in
5 methylene chloride (25 mL) at rt under argon was treated with DIEA (0.611 mL, 11 equiv). After 16 h the solvent was removed in vacuo and the residue dried in vacuo for 3 h. The residue was dissolved in methylene chloride (15 mL) and to this was added ether (30 mL) with slow stirring. The resulting suspension was left to stand overnight. The solid was collected by filtration,
10 washed with 2:1 ether/methylene chloride and dried in vacuo (411.5 mg, 68%). ¹H-NMR (CDCl₃) δ 1.30 (m, 4H), 1.45 (m, 4H), 1.62 (m, 2H), 1.89 (m, 2H), 2.10 (t, 4H), 2.58 (brt, 2H), 3.00 (brs, 4H), 3.06 (m, 4H), 3.31 (t, 2H), 3.73 (s, 6H), 4.38 (q, 2H), 4.68 (q, 2H), 5.21 (s, 4H), 6.53 (s, 2H), 6.77 (d, 4H), 7.03-7.38 (m, 34H), 7.41 (d, 8H), 7.57 (d, 4H), 8.22 (d, 4H), 8.79 (brs, 2H).
15 MS (ESI) 1893.9 (MH)⁺.

Example 27b**MEt-IDA-(Val-Lys(Mtr)-PABC-PNP)₂**

This was prepared from MEt-IDA-(Val-Lys(Mtr)-PABOH)₂ **26b** (300 mg, 0.205 mmol) as described above for **27a** (347.7 mg, 94%). ¹H-NMR
20 (CDCl₃) δ 0.88 (t, 12H), 1.25-1.75 (m, 12H), 1.93 (m, 2H), 2.07 (m, 4H), 2.71 (m, 2H), 3.21 (brs, 4H), 3.55 (m, 2H), 4.26 (m, 2H), 4.34 (m, 2H), 5.19 (s, 4H), 6.59 (s, 2H), 6.76 (d, 4H), 7.10-7.40 (m, 32H), 7.59 (m, 4H), 8.21 (d, 4H), 8.65 (br, 2H).
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Example 28a**MEt-IDA-(Phe-Lys(Mtr)-PABC-DOX)₂**

A solution of MEt-IDA-(Phe-Lys(Mtr)-PABC-PNP)₂ **27a** (410.3 mg, 0.217 mmol) and doxorubicin hydrochloride (283.0 mg, 2.25 equiv) in NMP (10 mL) at rt was treated with DIEA (0.094 mL, 2.5 equiv). After 48 h the mixture
30 was poured into ethyl acetate. The solution was washed with water (3x) and evaporated in vacuo. In each case some orange solid, that was shown by

TLC to contain product, was formed. This was collected by filtration and combined later with the organic-soluble components before chromatography. The combined materials were chromatographed on silica, eluting with 1) 6:4 , and 2) 1:1 methylene chloride/(5:1 ethyl

- 5 acetate/methanol), to give the product as an orange solid (351.8 mg, 60%).
 $^1\text{H-NMR}$ ($\text{CDCl}_3/\text{CH}_3\text{OD}$) δ 1.22 (brd, 10H), 1.40 (m, 4H), 1.51 (m, 4H), 1.75 (m, 6H), 2.07 (m, 4H), 2.20 (ABq, 4H), 2.50 (brt, 2H), 2.93 (brs, 4H), 2.99 (m, 4H), 3.23 (brs, 4H), 3.57 (m, 2H), 3.68 (s, 6H), 3.75 (m, 2H), 3.97 (s, 6H), 4.06 (m, 2H), 4.29 (m, 2H), 4.56 (m, 2H), 4.71 (s, 4H), 4.81 (ABq, 4H), 5.16
10 (brs, 2H), 5.42 (brs, 2H), 6.55 (s, 2H), 6.71 (d, 4H), 6.98-7.50 (m, 44H), 7.71 (t, 2H), 7.94 (d, 2H), 8.69 (br, 2H). MS (ESI) 2702.4 (MH) $^+$.

Example 28b

MEt-IDA-(Val-Lys(Mtr)-PABC-DOX) $_2$

- 15 This was prepared from MEt-IDA-(Val-Lys(Mtr)-PABC-PNP) $_2$ **27b** (6.351 g, 3.536 mmol) as described above for **28a** (2.79 g, 30%). $^1\text{H-NMR}$ ($\text{DMSO}-d_6$) δ 0.76 (ABq, 12H), 1.10 (d, 6H), 1.20-1.60 (m, 12H), 1.88 (m, 8H), 2.13 (q, 4H), 2.33 (m, 2H), 2.64 (m, 2H), 2.93 (m, 4H), 3.21 (brs, 4H), 3.64 (s, 6H), 3.70 (m, 2H), 3.91 (s, 6H), 4.15 (m, 4H), 4.26 (m, 2H), 4.55 (d, 4H), 4.70 (d,
20 2H), 4.88 (m, 4H), 5.19 (brs, 2H), 5.45 (s, 2H), 6.75 (d, 4H), 6.86 (s, 2H), 7.10 (m, 4H), 7.18 (m, 18H), 7.30 (d, 6H), 7.49 (m, 4H), 7.55 (m, 2H), 7.82 (m, 4H), 8.04 (m, 2H), 9.94 (br, 2H). MS (ESI) 2608.0 (MH) $^+$.

Example 29a

MEt-IDA-(Phe-Lys-PABC-DOX) $_2 \cdot 2\text{Cl}_2\text{CHCO}_2\text{H}$

- 25 A stirred solution of MEt-IDA-(Phe-Lys(Mtr)-PABC-DOX) $_2$ **28a** (192.0 mg, 0.071 mmol) in methylene chloride (15 mL), at rt was treated with anisole (1.54 mL, 200 equiv) and dichloroacetic acid (0.117 mL, 20 equiv). After 1h ethyl acetate (15 mL) was added. The resulting solid was collected by
30 filtration, washed with ethyl acetate and ether and dried in vacuo (181.0 mg, 100%). $^1\text{H-NMR}$ ($\text{CDCl}_3/\text{CH}_3\text{OD}$) δ 1.19 (d, 6H), 1.33 (m, 4H), 1.52 (m, 6H), 1.74 (m, 6H), 2.14 (ABq, 4H), 2.49 (brt, 2H), 2.93 (m, 12H), 3.25 (brs,

4H), 3.51 (m, 2H), 3.77 (m, 2H), 3.90 (s, 6H), 4.04 (m, 2H), 4.32 (m, 2H), 4.65 (s, 4H), 4.82 (m, 4H), 5.05 (brs, 2H), 5.33 (brs, 2H), 5.82 (brs, 2H), 6.63 (s, 2H), 7.01 (m, 14H), 7.32 (m, 6H), 7.63 (t, 2H), 7.79 (d, 2H). MS (ESI) 1078.8 (MH₂)²⁺, 2156.8 (MH)⁺.

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Example 29b

MEt-IDA-(Val-Lys-PABC-DOX)₂•2Cl₂CHCO₂H

This was prepared from MEt-IDA-(Val-Lys(Mtr)-PABC-DOX)₂ **28b** (232 mg, 0.0891 mmol) as described above for **29a** (171 mg, 78%). ¹H-NMR (DMSO-d₆) δ 0.78 (ABq, 12H), 1.10 (d, 6H), 1.15-1.70 (m, 12H), 1.78 (m, 2H), 1.92 (m, 2H), 2.11 (m, 4H), 2.72 (m, 4H), 2.94 (m, 4H), 3.68 (m, 2H), 3.94 (s, 6H), 4.15 (m, 4H), 4.26 (m, 2H), 4.55 (brs, 4H), 4.66 (brs, 2H), 4.86 (m, 4H), 5.19 (brs, 2H), 5.45 (s, 2H), 5.91 (s, 2H), 6.82 (d, 2H), 6.93 (s, 2H), 7.21 (d, 4H), 7.51 (d, 4H), 7.58 (m, 2H), 7.76 (m, 4H), 7.84 (m, 4H), 8.19 (m, 2H), 10.00 (brs, 6H). IR (KBr) 1648, 1709. MS (ESI) 1030.9 (MH₂)²⁺, 2060.8 (MH)⁺.

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Example 30

MEt-IDA-(Phe-Lys(Mtr)-PABC-MMC)₂

A stirred mixture of MEt-IDA-(Phe-Lys(Mtr)-PABC-PNP)₂ **27a** (183.8 mg, 0.0971 mmol), mitomycin C (73.1 mg, 2.25 equiv), N-hydroxybenzotriazole monohydrate (131.3 mg, 10 equiv) and freshly activated 4Å powdered sieves (0.5 g) in NMP (6 mL) under argon at rt was treated with DIEA (0.17 mL, 10 equiv). After 40 h the mixture was filtered and the filtrate diluted with ethyl acetate (80 mL). The solution was washed with water (4x), brine, dried over sodium sulfate and evaporated. The residue was chromatographed on silica, eluting with 2:1 methylene chloride/(5:1 ethyl acetate/methanol), to give the product as a light purple solid (201.7 mg, 91%). ¹H-NMR (CDCl₃) δ 1.27 (m, 4H), 1.42 (m, 4H), 1.60 (m, 2H), 1.69 (s, 6H), 1.83 (m, 2H), 2.06 (brt, 4H), 2.57 (m, 2H), 3.00 (m, 4H), 3.07 (brs, 4H), 3.13 (s, 6H), 3.28 (m, 4H), 3.41 (d, 2H), 3.59 (ABq, 2H), 3.72 (s, 6H), 4.20 (t, 2H), 4.41 (d and m, 3H), 4.66 (m, 2H), 4.88 (d, 4H), 5.10 (d, 2H), 5.39 (br, 6H), 6.49 (s, 2H), 6.72 (d,

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4H), 6.90-7.55 (m, 42H), 7.62 (br, 2H), 9.07 (br, 2H). MS (ESI) 1142.0 (MH₂)²⁺, 2282.0 (MH)⁺.

Example 31

5 **MEt-IDA-(Phe-Lys-PABC-MMC)₂•2Cl₂CHCO₂H**

A solution of MEt-IDA-(Phe-Lys(Mtr)-PABC-MMC)₂ **30** (134.6 mg, 0.0590 mmol) in methylene chloride (3 mL) at rt was treated with anisole (1.28 mL, 200 equiv) and 1M chloroacetic acid in methylene chloride (1.18 mL, 20 equiv). After 3.5 h ethyl acetate (30 mL) was added. the resulting purple solid was collected by filtration, washed with ethyl acetate and air-dried (97.8 mg, 82%). The product appeared to decompose in solution to one product by HPLC. ¹H-NMR (CDCl₃/CH₃OD) δ 1.33 (m, 4H), 1.63 (s and m, 12H), 1.77 (m, 2H), 2.52 (m, 2H), 2.82 (m, 6H), 3.03 (m, 2H), 3.10 (s and brs, 10H), 3.27 (m, 2H), 3.53 (ABq, 2H), 3.96 (brs, 4H), 4.11 (m, 2H), 4.35 (d and m, 3H), 4.53 (m, 2H), 4.78 (ABq, 2H), 4.94 (m, 4H), 6.62 (s, 2H), 7.10 (m, 10H), 7.19 (d, 4H), 7.41 (d, 4H). MS (FAB) 1738.3 (M+Na)⁺, 1776.4 (M+K)⁺.

Example 32

2'-Methoxytrityl-Paclitaxel

20 A stirred solution of paclitaxel (0.51 g, 0.597 mmol) and p-methoxytrityl chloride (4.63 g, 25 equiv) in methylene chloride (14 mL) under nitrogen at rt was treated with pyridine (1.23 mL, 25 equiv). After 16 h at rt the solvent was evaporated and the residue dissolved in ethyl acetate. The solution was washed with cold pH 5 buffer (2x100mL), water and brine, dried and
25 evaporated. The residue was chromatographed on silica, eluting with 3% methanol/methylene chloride, to give the product as a white solid (482 mg, 72%). ¹H NMR (CDCl₃) δ 1.11 (s, 3H), 1.17 (s, 3H), 1.55 (s, 3H), 1.67 (s, 3H), 1.90 and 2.54 (m, 2H), 2.26 (s, 3H), 2.51 (s, 3H), 2.54 (m, 2H), 3.66 (d, 1H), 3.78 (s, 3H), 4.21 (ABq, 2H), 4.41 (m, 1H), 4.63 (d, 1H), 4.92 (d, 1H),
30 5.62 (d, 1H), 5.70 (m, 2H), 6.22 (s, 1H), 6.74 (d, 2H), 7.09-7.60 (m, 23H), 7.80 (d, 2H), 8.09 (d, 2H). MS (FAB) 1148 (M+Na)⁺, 1164 (M+K)⁺.

Example 33**MEt-IDA-(Phe-Lys(Mtr)-PABC-7-Paclitaxel)₂**

A solution of paclitaxel-2'-Mtr **32** (789.3 mg, 0.701 mmol) in methylene chloride (5 mL) at 0°C under argon was treated with DIEA (0.122 mL, 1 equiv), pyridine (0.057 mL, 1 equiv) and diphosgene (0.043 mL, 0.5 equiv). The mixture was allowed to warm to rt for 1.5 h and then it was added, via syringe, to a stirred suspension of MEt-IDA-(Phe-Lys(Mtr)-PABOH)₂ **26a** (547.3 mg, 0.5 equiv) in methylene chloride (5 mL) and DIEA (0.122 mL, 1 equiv) at rt. After 16 h the mixture was diluted with ethyl acetate and washed with pH 5 buffer (0.5M biphthalate), water and brine, dried over sodium sulfate, and concentrated in vacuo. The residue was chromatographed on silica, eluting with 1) 5:1 and 2) 3:1 methylene chloride/(5:1 ethyl acetate/methanol), to give the product as a colorless glass (859.2 mg, 63%). Also recovered was a small amount of the mono-taxol adduct (172.0 mg, 18%). ¹H-NMR (CDCl₃/CH₃OD) δ 1.05-1.80 (4xm, 12H), 1.12 (s, 6H), 1.14 (s, 6H), 1.61 (s, 6H), 1.79 (s, 6H), 1.90 (m, 2H), 2.09 (m, 4H), 2.16 (s, 6H), 2.28 (s, 6H), 2.58 (m, 3H), 2.98 (brs, 4H), 3.05 (m, 4H), 3.37 (m, 2H), 3.75 (s, 6H), 3.78 (s, 6H), 4.16 (d, 2H), 4.27 (d, 2H), 4.32 (m, 2H), 4.63 (d and m, 4H), 4.92 (d, 2H), 5.16 (m, 4H), 5.49 (ABq, 2H), 5.62 (d, 2H), 5.71 (m, 4H), 6.33 (s, 2H), 6.53 (s, 2H), 6.75 (q, 8H), 6.97 (brd, 2H), 7.05-7.65 (m, 96H), 7.81 (d, 4H), 8.08 (d, 4H), 8.68 (brs, 2H).

Example 34**MEt-IDA-(Phe-Lys-PABC-7-Paclitaxel)₂•2ClCH₂CO₂H**

MEt-IDA-(Phe-Lys(Mtr)-PABC-7-Paclitaxel)₂ **33** (333.0 mg, 0.0861 mmol) in methylene chloride (3 mL) was treated with anisole (1.87 mL, 200 equiv) and 1M chloroacetic acid in methylene chloride (1.72 mL, 20 equiv). After 4 h the mixture was diluted with ether (30 mL). The resulting suspension was sonicated for several minutes and then the solid was collected by filtration and washed with ether (256.3 mg, 97%). The product was shown to decompose slowly in solution. ¹H-NMR (DMF-d₇) δ 1.15 (s, 6H), 1.19 (s, 6H), 1.77 (s, 6H), 1.94 (s, 6H), 1.30-2.00 (m, 12H), 2.15 (m, 4H), 2.59 (m, 4H), 3.09 (m, 8H), 3.93 (d, 2H), 4.16 (s, 4H), 4.69 (m, 4H), 4.90 (d, 2H), 5.07 (d,

2H), 5.16 (m, 4H), 5.56 (ABq, 2H), 5.67 (d, 2H), 5.70 (ABq, 2H), 6.14 (t, 2H), 6.42 (s, 2H), 6.49 (m, 2H), 7.02 (s, 2H), 7.10-7.90 (m, 44H), 8.11 (d, 4H), 9.09 (br, 2H). MS (ESI) 2778.2 (MH)⁺.

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Example 35**N-Carbobenzyloxy-N', N'-bis-(t-butoxycarbonylmethyl)propylenediamine**

This was prepared from N-carbobenzyloxypropylenediamine hydrochloride (3.4624 g, 14.15 mmol) as described above for **22** (5.3017 g, 86%). ¹H-NMR (CDCl₃) δ 1.45 (s, 18H), 1.63 (m, 2H), 2.72 (t, 2H), 3.32 (t, 2H), 3.38 (s, 4H), 5.11 (s, 2H), 6.26 (brt, 1H), 7.33 (m, 5H). MS (ESI) 437.5 (MH)⁺.

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Example 36**N, N-Bis-(t-butoxycarbonylmethyl)propylenediamine diacetic acid**

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A solution of N-carbobenzyloxy-N', N'-bis-(t-butoxycarbonylmethyl)propylenediamine **35** (5.3017 g, 12.144 mmol) in ethanol (75 mL) was filtered through a bed of Raney nickel into a solution of acetic acid (2.1 mL, 3 equiv) in ethanol (75 mL). The filtrate was degassed with nitrogen and treated with 10% palladized charcoal (0.4 g). The mixture was hydrogenated at 50 psi on a Parr apparatus for 16 h and then the catalyst was filtered off on a bed of celite. The filtrate was concentrated in vacuo and the residue flushed with chloroform (150 mL). The crude product was carried on without further purification.

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Example 37**N-Maleoyl-N', N'-bis-(t-butoxycarbonylmethyl)propylenediamine**

This was prepared from N, N-bis-(t-butoxycarbonylmethyl)propylenediamine diacetic acid **36** (12.14 mmol) as described above for **24**. The crude product was chromatographed on silica, eluting with 3:1 hexane/ethyl acetate, to give the product as a pale-yellow oil (2.0653 g, 44%).

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(3 steps)). $^1\text{H-NMR}$ (CDCl_3) δ 1.42 (s, 18H), 1.74 (m, 2H), 2.72 (t, 2H), 3.40 (s, 4H), 3.60 (t, 2H), 6.68 (s, 2H). MS (DCI) 383 (MH) $^+$, 327 (MH-C₄H₉) $^+$.

Example 38

5 **N-Maleoyl-N', N'-bis-(carboxymethyl)propylenediamine• p-toluenesulfonic acid**

A stirred solution of N-maleoyl-N', N'-bis-(t-butoxycarbonylmethyl)-propylenediamine **37** (2.06 g, 5.39 mmol) in methylene chloride (25 mL) at rt was treated with p-toluenesulfonic acid monohydrate (3.07 g, 3 equiv). After
10 16 h the solvent was removed in vacuo. The residue was triturated with ether four times (each time the ether was decanted off). The resulting semi-solid material was flushed with chloroform (2 x 150 mL). NMR showed 1.4-1.5 moles of p-toluenesulfonic acid per mole of product (2.68 g, 97%). $^1\text{H-NMR}$ (CD_3OD) δ 1.96 (m, 2H), 2.29 (s, 4.2H), 3.31 (m, 2H), 3.51 (t, 2H), 4.13
15 (s, 4H), 6.78 (s, 2H), 7.15 (d, 2.8H), 7.62 (d, 2.8H).

Example 39

MPr-IDA-(Phe-Lys(Mtr)-PABOH)₂

This was prepared from N-maleoyl-N', N'-bis-(carboxymethyl)propylenediamine-p-toluenesulfonic acid **38** (2.6842 g, 5.25
20 mmol) and Phe-Lys(Mtr)-PABOH **10a** (7.0432 g, 2 equiv) as described above for **26a**, except that the crude product was chromatographed on silica, eluting with 1) 2:1 and 2) 1.5:1 methylene chloride/(5:1 ethyl acetate/methanol), to give the product as a white solid (4.5253 g, 55%). $^1\text{H-NMR}$ ($\text{CDCl}_3/\text{CD}_3\text{OD}$) δ 1.24 (m, 4H), 1.41 (m, 4H), 1.52 (m, 2H), 1.83 (m, 2H), 2.06 (brt, 4H), 2.23 (brt, 2H), 2.88 (ABq, 4H), 3.09 (m, 4H), 3.27 (t, 2H), 3.73 (s, 6H), 4.36 (q, 2H), 4.53 (brs, 4H), 4.71 (q, 2H), 6.61 (s, 2H), 6.75 (d, 4H), 7.05-7.48 (m, 42H), 8.74 (brs, 2H). MS (ESI) 1576.4 (MH) $^+$, 1303.6 (M-Mtr) $^+$.

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Example 40**MPr-IDA-(Phe-Lys(Mtr)-PABC-PNP)₂**

This was prepared from MPr-IDA-(Phe-Lys(Mtr)-PABOH)₂ **39** (1.7993 g, 1.143 mmol) as described above for **27a** (1.8909 g, 87%). ¹H-NMR (CDCl₃) δ 1.29 (m, 4H), 1.42 (m, 4H), 1.60 (m, 2H), 1.88 (m, 2H), 2.07 (brt, 4H), 2.32 (brt, 2H), 2.92 (ABq, 4H), 3.08 (m, 4H), 3.29 (t, 2H), 3.72 (s, 6H), 4.35 (q, 2H), 4.66 (q, 2H), 5.20 (s, 4H), 6.63 (s, 2H), 6.75 (d, 4H), 7.05-7.45 (m, 42H), 7.52 (d, 4H), 8.21 (d, 4H), 8.71 (brs, 2H). MS (ESI) 1906.0 (MH)⁺, 1634.4 (M-Mtr)⁺. MS (ESI) 1907.8 (MH)⁺.

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Example 41**MPr-IDA-(Phe-Lys(Mtr)-PABC-DOX)₂**

This was prepared from MPr-IDA-(Phe-Lys(Mtr)-PABC-PNP)₂ **40** (722.7 mg, 0.379 mmol) as described above for **28a**. The crude product was chromatographed on silica, eluting with 1) 20:1 and 2) 15:1 methylene chloride/methanol, to give the product as an orange solid (869.4 mg, 84%). ¹H-NMR (CDCl₃/CD₃OD) δ 1.16 (d, 10H), 1.42 (m, 4H), 1.49 (m, 4H), 1.82 (m, 6H), 2.00 (m, 2H), 2.12 (m, 4H), 2.19 (m, 4H), 2.53 (brt, 2H), 2.89 (brs, 4H), 2.99 (m, 4H), 3.25 (brs, 4H), 3.49 (m, 2H), 3.67 (s, 6H), 3.73 (m, 2H), 3.99 (s, 6H), 4.12 (m, 2H), 4.31 (m, 2H), 4.55 (m, 2H), 4.71 (s, 4H), 4.92 (ABq, 4H), 5.15 (brs, 2H), 5.40 (brs, 2H), 6.55 (s, 2H), 6.69 (d, 4H), 7.00-7.50 (m, 44H), 7.70 (t, 2H), 7.96 (d, 2H), 8.71 (br, 2H). MS (ESI) 2716.8 (MH)⁺.

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Example 42**MPr-IDA-(Phe-Lys-PABC-DOX)₂ • 2Cl₂CHCO₂H**

This was prepared from MPr-IDA-(Phe-Lys(Mtr)-PABC-DOX)₂ **41** (714.3 mg, 0.263 mmol) as described above for **29a** (645.9 mg, 96%). ¹H-NMR (DMF-d₇) δ 1.23 (d, 6H), 1.54 (m, 6H), 1.73 (m, 6H), 2.02 (m, 2H), 2.26 (ABq, 2H), 2.48 (m, 4H), 3.09 (m, 4H), 3.12 (brs, 4H), 3.29 (m, 6H), 3.91 (m, 2H), 4.10 (s, 6H), 4.32 (q, 2H), 4.56 (m, 2H), 4.80 (s and m, 6H), 5.13 (brs, 2H), 5.41 (brs, 2H), 5.68 (br, 2H), 6.17 (s, 2H), 6.72 (d, 2H), 6.99 (s, 2H), 7.27 (m, 18H), 7.70

30

(m, 4H), 7.92 (m, 2H), 8.52 (d, 2H), 8.69 (d, 2H), 10.22 (brs, 2H). MS-(ESI) —
2171.3 (MH)⁺.

Example 43

5 **Di-t-butyliminodiacetate**

A stirred mixture of glycine t-butyl ester hydrochloride (5.0811 g, 30.31 mmol) and potassium bicarbonate (6.3730 g, 2.1 equiv) in DMF (40 mL) at 0°C was treated with t-butyl bromoacetate (4.9 mL, 1 equiv). The ice bath was removed and the mixture was allowed to stir at rt for 14 h. The solvent
10 was removed in vacuo at 30°C and the residue was partitioned between ethyl acetate and 50% sat. sodium bicarbonate. The organic phase was washed with water (3x) and brine, dried over sodium sulfate, and concentrated in vacuo. The residue was chromatographed on silica, eluting with 1) 5% and 2) 25% ethyl acetate/hexane, to give the product as a
15 colorless oil (4.9310 g, 66%). ¹H-NMR (CDCl₃) δ 1.46 (s, 18H), 3.55 (s, 4H). MS (DCI) 246 (MH)⁺, 190 (MH-C₄H₉)⁺.

Example 44

20 **N-Maleoyl-N', N'-bis-(t-butoxycarbonyl-methyl)propionamide**

A suspension of 2-maleimidopropionic acid (1.0007 g, 5.916 mmol) in methylene chloride (12 mL) at rt was treated with oxalyl chloride (1.2 mL, 2.1 equiv) and DMF (1 drop). After 4 h the solvents were evaporated. The residue was flushed with dry methylene chloride (2 x 20 mL) and dissolved
25 in methylene chloride (2 mL). To this was added a solution of di-t-butyliminodiacetate **43** (1.4510 g, 1 equiv) and DIEA (1.24 mL, 1.2 equiv) in methylene chloride (6 mL), dropwise over 15 min. After 14 h the mixture was diluted with ethyl acetate and washed with 15% citric acid, water and brine. The organic phase was dried over sodium sulfate and concentrated in vacuo
30 to give a pale-yellow glass which was pure enough by NMR to carry to the next step (2.3448 g, 100%). ¹H-NMR (CDCl₃) δ 1.43 (s, 9H), 1.46 (s, 9H),

2.64 (t, 2H), 3.88 (t, 2H), 3.97 (s, 2H), 4.04 (s, 2H), 6.69 (s, 2H). MS_x(DCI) —
397 (MH)⁺, 341 (MH-C₄H₉)⁺.

Example 45

5 **N-Maleoyl-N', N'-bis-(carboxymethyl)- propionamide**

A stirred mixture of N-maleoyl-N', N'-bis-(t-butoxycarbonylmethyl)propionamide **44** (1.9100 g, 4.82 mmol) in methylene chloride (40 mL) at rt was treated with trifluoroacetic acid (20 mL). The mixture was sonicated at rt for 20 min and then left to stand for 3.5 h. The
10 reaction was concentrated in vacuo and the residue flushed with chloroform (2x50 mL) and then triturated with ether. The resulting pale-yellow solid was collected by filtration and washed with ether (1.2273 g, 90%). ¹H-NMR (CD₃OD/CDCl₃) δ 2.61 (t, 2H), 3.69 (t, 2H), 4.02 (s, 2H), 4.13 (s, 2H), 6.71 (s, 2H). MS (FAB⁻) 283.1 (M-H)⁻. HRMS Calcd: 285.0723. Found: 285.0712.

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Example 46

MP-IDA-(Phe-Lys(Mtr)-PABOH)₂

This was prepared from N-maleoyl-N', N'-bis-(carboxymethyl)propionamide **45** (142.1 mg, 0.500 mmol) and Phe-Lys(Mtr)-PABOH **10a** (671.0 mg, 2
20 equiv) as described above for **26a**, except that the crude product was chromatographed on silica, eluting with 1) 2:1 and 1.5:1 methylene chloride/(5:1 ethyl acetate/methanol), to give the product as a white solid (349.7 mg, 44%). ¹H-NMR (CDCl₃/CD₃OD) δ 1.00-1.90 (m, 12H), 2.07 (brt, 4H), 2.27 (brt, 2H), 3.06 (m, 4H), 3.32-3.82 (m, 6H), 3.70 (s, 6H), 4.26 (m,
25 1H), 4.35 (m, 2H), 4.52 (d, 4H), 4.62 (m, 1H), 6.51 (s, 2H), 6.72 (q, 4H), 7.00-7.52 (m, 42H). MS (FAB) 1612.5 (M+Na)⁺, 1628 (M+K)⁺.

Example 47

MP-IDA-(Phe-Lys(Mtr)-PABC-PNP)₂

30 This was prepared from MP-IDA-(Phe-Lys(Mtr)-PABOH)₂ **46** (225.4 mg, 0.142 mmol) as described above for **27a** (195.5 mg, 72%). ¹H-NMR (CDCl₃/CD₃OD) δ 1.05-1.95 (m, 12H), 2.05 (brt, 4H), 2.33 (m, 2H), 3.08 (m,

4H), 3.55 (t, 2H), 3.70 (s, 6H), 3.60-3.99 (m, 4H), 4.52 (m, 2H), 4.63 (m, 2H), 5.21 (d, 4H), 6.57 (s, 2H), 6.74 (q, 4H), 7.00-7.65 (m, 46H), 8.20 (q, 4H). MS (ESI) 1920.8 (MH)⁺, 1647.8 (MH-Mtr)⁺.

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Example 48**MP-IDA-(Phe-Lys(Mtr)-PABC-DOX)₂**

This was prepared from MP-IDA-(Phe-Lys(Mtr)-PABC-PNP)₂ **47** (195.0 mg, 0.102 mmol) as described above for **28a**. The crude product was chromatographed on silica, eluting with 1) 20:1 and 2) 15:1 methylene chloride/methanol, to give the product as an orange solid (117.7 mg, 42%).
¹H-NMR (CDCl₃/CD₃OD) δ 1.21 (m and d, 10H), 1.42 (m, 4H), 1.59 (m, 2H), 1.80 (m, 4H), 2.06 (m, 6H), 3.02 (m, 8H), 3.40-3.90 (m, 6H), 3.69 (s, 6H), 4.00 (s, 6H), 4.07 (m, 2H), 4.29 (m, 2H), 4.48 (m, 4H), 4.72 (s, 4H), 4.90 (m, 4H), 5.19 (brs, 2H), 5.42 (brs, 2H), 6.51 (s, 2H), 6.71 (m, 4H), 6.90-7.55 (m, 44H), 7.70 (t, 2H), 7.92 (m, 2H). MS (ESI) 1365.4 (MH₂)⁺², 2728.8 (MH)⁺.

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Example 49**MP-IDA-(Phe-Lys-PABC-DOX)₂•2Cl₂CHCO₂H**

This was prepared from MP-IDA-(Phe-Lys(Mtr)-PABC-DOX)₂ **48** (91.0 mg, 0.033 mmol) as described above for **29a** (76.4 mg, 94%).
¹H-NMR (CDCl₃/CD₃OD) δ 1.21 (d, 6H), 1.30-1.182 (m, 14H), 2.01 (t, 2H), 2.30 (m, 4H), 2.53 (m, 2H), 3.04 (m, 12H), 3.50 (m, 2H), 3.72 (m, 2H), 3.92 (s, 6H), 4.03 (m, 2H), 4.19 (m, 2H), 4.52 (m, 2H), 4.65 (s, 4H), 4.84 (m, 4H), 5.12 (brs, 2H), 5.37 (brs, 2H), 5.79 (s, 2H), 6.57 (s, 2H), 6.73 (d, 2H), 6.95-7.50 (m, 20H), 7.68 (t, 2H), 7.89 (m, 2H). MS (ESI) 1092.4 (MH₂)⁺², 2185.6 (MH)⁺.

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Example 50**Z-βAla**

A stirred solution of β-alanine (50.10 g, 0.562 mol) in 2M NaOH (281 mL) at 0°C was treated with benzyl chloroformate (88 mL, 1.1 equiv), dropwise over 1 h. Stirring was continued for 1.5 h at 0°C, followed by 1.5 h at rt. The resulting mixture was extracted with ether (4x500 mL). The aqueous phase

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was acidified with 5M HCl (120 mL). The resulting white precipitate was collected by filtration, washed with water and dissolved in methylene chloride. The solution was dried over sodium sulfate and concentrated in vacuo to give a white solid (114.95 g, 92%). ¹H-NMR (CDCl₃) δ 2.61 (t, 2H), 3.48 (q, 2H), 5.12 (s, 2H), 5.29 (br, 1H), 7.37 (s, 5H). MS (DCI) 224 (MH)⁺.

Example 51

Z-βAla-OSu

10 A mixture of Z-βAla **50** (37.42 g, 0.168 mol) and N-hydroxysuccinimide (19.34 g, 1 equiv) in methylene chloride (700 mL) was stirred at 0°C and treated with DCC (34.69 g, 1 equiv). The reaction was allowed to warm to rt. After 16 h the DCU was removed by filtration and the filtrate was concentrated in vacuo to give a white foam (53.80 g, 100%). ¹H-NMR
15 (CDCl₃) δ 2.86 (s and m, 6H), 3.60 (q, 2H), 5.12 (s, 2H), 5.41 (m, 1H), 7.37 (m, 5H). MS (DCI) 321 (MH)⁺.

Example 52

Z-βAla-IDA

20 A stirred solution of iminodiacetic acid (16.84 g, 84.32 mmol) and lithium hydroxide monohydrate (10.62 g, 2 equiv) in water (240 mL) at 0°C was treated with a solution of Z-βAla-OSu **51** (27.01 g, 0.667 equiv) in DME (130 mL). After 15 min the ice bath was removed and the reaction was stirred at rt for 16 h. The mixture was concentrated to a volume of 200 mL on the rotary
25 evaporator. The resulting suspension was extracted with ethyl acetate (2x250 mL), and then the aqueous phase was acidified to pH 1-2 with 85% phosphoric acid. The mixture was extracted with 10% isopropanol/ethyl acetate (3x200 mL). The combined organic phases were washed with 2% phosphoric acid and brine, dried over sodium sulfate, and evaporated to
30 give a colorless glass which was flushed with chloroform, giving a foam (21.36 g, 75%). ¹H-NMR (CD₃OD) δ 2.51 (t, 2H), 3.32 (t, 2H), 4.04 (s, 2H), 4.15 (s, 2H), 4.98 (s, 2H), 7.25 (m, 5H). MS (FAB) 339 (MH)⁺, 361 (M+Na)⁺,

377 (M+K)⁺. Anal. Calcd for C₁₅H₁₈N₂O₇·H₂O: C-50.56, H-5.66, N-7.86. —
Found: C-50.56, H-5.45, N-8.21.

Example 53

5 **Z-βAla-IDA-(OSu)₂**

This was prepared from Z-βAla-IDA **52** (21.36 g, 63.12 mmol) as described above for **51**. The crude product was used without purification. ¹H-NMR (CDCl₃/CD₃OD) δ 2.62 (t, 2H), 2.83 (s, 8H), 3.53 (m, 2H), 4.56 (s, 4H), 5.08 (s, 2H), 5.58 (brt, 1H), 7.35 (m, 5H). Anal. Calcd for C₂₃H₂₄N₄O₁₁·0.5H₂O: 10 C-51.02, H-4.65, N-10.35. Found: C-50.83, H-5.13, N-10.75.

Example 54

Z-βAla-IDA-(βAla-O-t-Bu)₂

A stirred suspension of Z-βAla-IDA-(OSu)₂ **53** (ca. 63.12 mmol) and β- 15 alanine t-butyl ester hydrochloride (25.80 g, 2.25 equiv) in DME (250 mL) at rt were treated with DIEA (24.7 mL, 2.25 equiv). After 12 h the solvents were evaporated and the residue partitioned between ethyl acetate (400 mL) and 2% phosphoric acid. The organic phase was washed with water and brine, dried over sodium sulfate and evaporated. The residue was 20 chromatographed on silica, eluting with 1) ethyl acetate, 2) 15:1, and 3) 10:1 ethyl acetate/methanol, to give the product as a thick, colorless oil (26.56 g, 71%). ¹H-NMR (CDCl₃) δ 1.41 (s, 18H), 2.46 (m, 6H), 3.49 (m, 6H), 3.88 (s, 2H), 3.97 (s, 2H), 5.09 (s, 2H), 5.56 (brt, 1H), 6.89 (brt, 1H), 7.33 (m, 5H), 8.74 (brt, 1H). MS (FAB) 593 (MH)⁺, 615 (M+Na)⁺, 631 (M+K)⁺. Anal. Calcd for 25 C₂₉H₄₄N₄O₉·H₂O: C-57.04, H-7.59, N-9.17. Found: C-57.15, H-7.36, N-9.46.

Example 55

βAla-IDA-(βAla-O-t-Bu)₂

30 Z-βAla-IDA-(O-t-Bu)₂ **54** (16.93 g, 28.56 mmol) in absolute ethanol (100 mL) was degassed by bubbling nitrogen through it for 30 min. Palladized charcoal (10%, 1 g) was carefully added and the mixture was hydrogenated

on a Parr apparatus at 50 psi for 16 h. The catalyst was then filtered off on a bed of celite and the filtrate concentrated in vacuo. The residue was flushed with methylene chloride, giving a colorless foam (13.11 g, 100%). $^1\text{H-NMR}$ (CDCl_3) δ 1.47 (s, 18H), 2.45 (q, 6H), 3.01 (t, 2H), 3.51 (m, 4H), 3.91 (s, 2H), 4.04 (s, 2H), 6.98 (brt, 1H), 8.93 (brt, 1H). MS (FAB) 459 (MH) $^+$, 481 (M+Na) $^+$, 497 (M+K) $^+$. Anal. Calcd for $\text{C}_{21}\text{H}_{38}\text{N}_4\text{O}_7 \cdot 2.5\text{H}_2\text{O}$: C-50.09, H-8.61, N-11.12. Found: C-50.17, H-8.61, N-10.51.

Example 56

10 **Maleoyl- β Ala-IDA-(β Ala-O-t-Bu) $_2$**

A stirred solution of β Ala-IDA-(O-t-Bu) $_2$ **55** (6.20g, 13.5 mmol) in chloroform (100 mL) at rt was treated with maleic anhydride (1.46 g, 1.1 equiv). After 1.5 h the mixture was concentrated in vacuo. The residue was flushed with hexane and carried on without further purification. $^1\text{H-NMR}$ (CDCl_3) δ 1.50 (s, 18H), 2.48 (m, 4H), 2.57 (m, 2H), 3.48 (m, 4H), 3.67 (q, 2H), 3.92 (s, 2H), 4.06 (s, 2H), 6.38 (ABq, 2H), 7.39 (brt, 1H), 7.91 (brt, 1H), 8.22 (brt, 1H). MS (FAB) 459.1 (MH) $^+$, 481.2 (M+Na) $^+$, 497.1 (M+K) $^+$.

Example 57

20 **MP-IDA-(β Ala-O-t-Bu) $_2$**

A solution of maleoyl- β Ala-IDA-(β Ala-O-t-Bu) $_2$ **56** (7.50 g, 13.5 mmol) in acetonitrile (150 mL) under argon at 0°C was treated with trimethylsilyl chloride (8.6 mL, 5.02 equiv) followed by triethylamine (9.4 mL, 5.02 equiv). The mixture was heated at reflux for 4 h, cooled to rt, and then stored overnight at -20°C. The resulting solid triethylamine hydrochloride was filtered off and the filtrate concentrated in vacuo. The residue was partitioned between ethyl acetate and water. The aqueous phase was extracted with more ethyl acetate. The combined organic phases were dried over sodium sulfate and evaporated. The residue was chromatographed on silica, eluting with 1) 5%, 2) 7%, and 3) 10% methanol/methylene chloride, to give the product as a pale-yellow gum (4.83 g, 66%). $^1\text{H-NMR}$ (CDCl_3) δ 1.42 (s, 18H), 2.47 (m, 4H), 2.61 (t, 2H), 3.50 (m, 4H), 3.81 (t, 2H), 3.88 (s,

2H), 3.99 (s, 2H), 6.60 (brt, 1H), 6.69 (s, 2H), 8.82 (brt, 1H). MS (FAB) 561.2 (M+Na)⁺, 577 (M+K)⁺. Anal. Calcd for C₂₅H₃₈N₄O₇·1.5H₂O: C-53.09, H-7.30, N-9.91. Found: C-53.13, H-6.98, N-10.05.

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Example 58

MP-IDA-(βAla)₂

A solution of MP-IDA-(βAla-O-t-Bu)₂ **57** (3.50 g, 6.50 mmol) in methylene chloride (40 mL) at rt was treated with trifluoroacetic acid (30 mL). the mixture was sonicated for 10 min and then left to stand for 1 h. The solvents were evaporated on the rotovap and the residue flushed with chloroform (3x) and ether (4x). The crude product was carried on without further purification. ¹H-NMR (CD₃OD) δ 2.47 (m, 4H), 2.53 (m, 2H), 3.40 (m, 4H), 3.69 (t, 2H), 3.89 (s, 2H), 4.04 (s, 2H), 6.71 (s, 2H). MS (FAB) 433.0 (M+Li)⁺. Anal. Calcd for C₁₇H₂₂N₄O₉·3H₂O: C-42.50, H-5.87, N-11.66. Found: C-42.44, H-5.92, N-11.54.

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Example 59

MP-IDA-(βAla-OSu)₂

A stirred solution of MP-IDA-(βAla)₂ **58** (136.0 mg, 0.319 mmol) and NHS (73.5 mg, 2 equiv) in DMF (2 mL) at rt was treated with DCC (265.0 mg, 4.01 equiv). The mixture was stirred at rt for 16 h and then was carried on directly to the next step without isolation of the product.

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Example 60

MP-IDA-(βAla-Phe-Lys(Mtr)-PABOH)₂

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The crude reaction mixture containing MP-IDA-(βAla)₂ **58** (ca. 0.319 mmol) was diluted with DMF (5 mL) and then treated with Phe-Lys(Mtr)-PABOH **10a** (429.1 mg, 2.0 equiv). After stirring at rt for 5 h the solvents were removed under high vacuum and the residue was triturated with ether. The resulting solid was collected by filtration and then chromatographed on silica, eluting with 5% methanol/ethyl acetate to remove impurities, and 10% methanol/methylene chloride to elute the product (220.1 mg, 40%). ¹H-NMR

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(DMSO-d₆) δ 1.15-1.75 (m, 12H), 1.92 (m, 4H), 2.21 (m, 4H), 2.41 (t, 2H), 2.72 (m, 2H), 3.01 (m, 2H), 3.13 (m, 4H), 3.58 (m, 2H), 3.67 (s, 6H), 3.78 (d, 2H), 3.90 (brs, 2H), 4.38 (m, 2H), 4.42 (d, 4H), 4.53 (m, 2H), 5.09 (t, 2H), 7.80 (d, 4H), 6.96 (s, 2H), 7.05-7.30 (m, 32H), 7.38 (d, 6H), 7.52 (d, 4H), 8.08 (m, 2H), 8.13 (d, 2H), 9.91 (brs, 2H). MS (FAB) 1754.0 (M+Na)⁺, 1770.0 (M+K)⁺.

Example 61

MP-IDA-(β Ala-Phe-Lys(Mtr)-PABC-PNP)₂

A stirred solution of MP-IDA-(β Ala-Phe-Lys(Mtr)-PABOH)₂ **60** (95 mg, 0.0548 mmol) in 1:1 DMF/methylene chloride (6 mL) at rt was treated with bis-p-nitrophenylcarbonate (100 mg, 6 equiv) and DIEA (0.057 mL, 6.0 equiv). After 16 h the solvents were removed under high vacuum and the residue was triturated with 1:1 methylene chloride/ether until it became a solid. This suspension was stored in the freezer for 1.5 h and then the solid was collected by filtration and washed with ether (105 mg, 95%). ¹H-NMR (DMSO-d₆) δ 1.20-1.75 (m, 12H), 1.93 (m, 4H), 2.21 (m, 4H), 2.41 (m, 2H), 2.71 (t, 2H), 3.01 (d, 2H), 3.14 (m, 4H), 3.56 (m, 2H), 3.67 (s, 6H), 3.78 (brs, 2H), 3.91 (brs, 2H), 4.37 (m, 2H), 4.56 (m, 2H), 5.23 (s, 4H), 6.80 (d, 4H), 6.94 (s, 2H), 7.05-7.50 (m, 38H), 7.56 (d, 4H), 7.64 (d, 4H), 8.09 (m, 2H), 8.18 (d, 2H), 8.30 (d, 4H), 10.08 (brs, 2H). MS (ESI) 2062.8 (MH)⁺.

Example 62

MP-IDA-(β Ala-Phe-Lys(Mtr)-PABC-DOX)₂

MP-IDA-(β Ala-Phe-Lys(Mtr)-PABC-PNP)₂ **61** (1.445 g, 0.701 mmol) and doxorubicin hydrochloride (1.010 g, 2.48 equiv) in DMF (55 mL) were treated with DIEA (0.31 mL, 2.50 equiv). After 40 h at rt the solvent was evaporated under high vacuum and the residue was triturated with 1:1 methylene chloride/ether. The resulting suspension was sonicated briefly and then the solid was collected by filtration and washed with ether. The solid was dissolved as far as possible in 10% methanol/chloroform (100 mL). The mixture was filtered and the filtrate was concentrated on the rotovap to ca. 50 mL. Chloroform (50 mL) was added and the procedure was repeated two more times. The resulting solid was collected by filtration and washed with

chloroform and then ether (1.22 g, 61%). $^1\text{H-NMR}$ (DMSO- d_6) δ 1.13 (d, 6H), 1.20-1.75 (m, 12H), 1.81 (m, 2H), 1.92 (m, 4H), 2.19 (m, 6H), 2.30 (m, 2H), 2.72 (m, 2H), 2.98 (m, 6H), 3.12 (m, 2H), 3.45 (m, 2H), 3.56 (t, 2H), 3.69 (s, 6H), 3.73 (m, 2H), 3.89 (m, 2H), 3.96 (s, 6H), 4.14 (m, 2H), 4.33 (m, 2H), 4.57 (d and m, 6H), 4.70 (d, 2H), 4.84 (t, 2H), 4.91 (m, 4H), 5.22 (brs, 2H), 5.46 (s, 2H), 6.79 (d, 4H), 6.93 (s, 2H), 7.05-7.30 (m, 32H), 7.33 (d, 6H), 7.52 (d, 4H), 7.60 (m, 2H), 7.88 (d, 2H), 8.06 (m, 2H), 8.13 (d, 2H), 9.96 (brs, 2H). MS (ESI) 2872.6 (MH) $^+$.

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Example 63

MP-IDA-(β Ala-Phe-Lys-PABC-DOX) $_2$ •2Cl $_2$ CHCO $_2$ H A stirred suspension of MP-IDA-(β Ala-Phe-Lys(Mtr)-PABC-DOX) $_2$ **62** (132 mg, 0.0459 mmol) in methylene chloride (8 mL) was treated with anisole (1 mL, 200 equiv) and dichloroacetic acid (0.076 mL, 20 equiv). The mixture immediately became homogeneous. After 5 min an orange precipitate began to form. After 2 h ethyl acetate (25 mL) was added and the suspension stirred for 30 min. The solid was collected by filtration and washed with ethyl acetate and then ether (89 mg, 80%). $^1\text{H-NMR}$ (DMSO- d_6) δ 1.12 (d, 6H), 1.22-1.90 (14H), 2.20 (m, 8H), 2.74 (m, 6H), 2.97 (m, 6H), 3.15 (m, 2H), 3.43 (brs, 2H), 3.56 (m, 2H), 3.72 (m, 2H), 3.97 (s, 6H), 4.14 (m, 2H), 4.38 (m, 2H), 4.56 (s and m, 6H), 4.70 (m, 2H), 4.89 (m, 4H), 5.21 (brs, 2H), 5.46 (brs, 2H), 6.01 (s, 2H), 6.81 (d, 2H), 6.97 (s, 2H), 7.21 (m, 14H), 7.56 (d, 4H), 7.61 (m, 2H), 7.77 (m, 2H), 7.89 (d, 2H), 8.12 (m, 2H), 10.01 (m, 2H). MS (ESI) 2325.6 (MH) $^+$.

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Example 64**M-Et-IDA-(Phe-Lys-PABC-DOX= β -Ala-mTEG) $_2$**

MTEG- β -Ala Hydrazide (254 mg, 866 μ mole) was dissolved in 10 ml methanol along with 50 μ l TFA. This solution was added to a suspension of M-Et-IDA-(Phe-Lys-PABC-DOX) $_2$ (276 mg, 108 μ mole) in 40 ml anhydrous methanol. The reaction was stirred for 3.5 hr. at room temperature. The reaction mixture was rotary evaporated to the point of precipitation, then 1 ml

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CH₂Cl₂ was added to redissolve. This was added dropwise to 250 ml ether, precipitating a red solid. The solid was filtered, washed with ether, and dried under high vacuum to yield M-Et-IDA-(Phe-Lys-PABC-DOX=β-Ala-mTEG)₂ (300 mg, 90%).

- 5 ¹H-NMR (d⁷-DMF): (selected peaks) δ 1.23 (d, 6H), 2.34 (t, 4H), 3.28 (s, 6H), 3.55 (s, 24H), 4.03 (s, 6H), 4.10 (t, 4H), 6.98 (s, 2H).

Mass Spec.: FAB 1354.6 (M+2H)²⁺

Elemental Analysis for C₁₃₂H₁₆₄N₁₈O₄₄ •4.0H₂O•3.0TFA: Theoretical C, 53.11; H, 5.65; N, 8.08. Found C, 52.99; H, 5.96; N, 8.64.

- 10 FTIR: 3412, 2938, 1702, 1681, 1616, 1520, 1412, 832, 696 cm⁻¹.

Example 65

M-Et-IDA-(Phe-Lys-PABC-DOX=β-Ala)₂

- β-Ala Hydrazide trifluoroacetate (578 mg, 1.75 mmole) was dissolved in 10 ml methanol along with 50 ul TFA. This solution was added to a suspension of M-Et-IDA-(Phe-Lys-PABC-DOX)₂ (111 mg, 43.6 umole) in 15 ml anhydrous methanol. The reaction was stirred for 1.5 hr. at room temperature. The reaction mixture was rotary evaporated to a volume of 3 ml, then added dropwise to 200 ml acetonitrile, precipitating a red solid. The solid was filtered, washed with acetonitrile, and dried under high vacuum to yield M-Et-IDA-(Phe-Lys-PABC-DOX=β-Ala)₂ (117 mg, 99%).

¹H-NMR (d⁶-DMSO): (selected peaks) δ 1.13 (d, 6H), 3.95 (s, 6H), 6.96 (s, 2H).

- 25 **Example 66**

N-t-Butoxycarbonyl-3-aminopropanol

- A solution of 3-aminopropanol (15 mL, 0.20 mol) in methylene chloride (50 mL) at rt was treated with t-butylpyrocarbonate (42.02 g, 1 equiv) in methylene chloride (25 mL), dropwise over 2 h. After stirring overnight the mixture was evaporated, flushed with heptane (3x), and dried in vacuo to give the product as a thick oil (34.31 g, 100%). ¹H NMR δ 1.48 (s, 9H), 3.12 (br, 1H), 3.22 (q, J=7.5 Hz, 2H), 3.60 (q, J=7.8 Hz, 2H), 4.81 (br, 1H). MS

(DCI) 176 (MH)⁺. Anal calcd for C₈H₁₇NO₃: C-54.84, H-9.78, N-7.99. Found: C-54.80, H-9.82, N-7.97.

Example 67

5 **N-t-Butoxycarbonyl-3-aminopropyl mesylate**

The protected amino alcohol **64** (6.601 g, 37.67 mmol) in methylene chloride (45 mL) at 0°C under argon was treated with methanesulfonyl chloride (2.92 mL, 1.0 equiv) and triethylamine (5.25 mL, 1.0 equiv), dropwise over 30 min. The mixture was stirred for 16 h and then the solvent
10 was evaporated. The residue was partitioned between ether (200 mL) and water (250 mL) and the organic phase was washed with 10% citric acid, water and brine, dried over sodium sulfate and evaporated. The residue was chromatographed on silica, eluting with 2% methanol/methylene chloride. The product was found to be unstable over extended periods at rt
15 (5.800 g, 61%). ¹H NMR δ 1.34 (s, 9H), 1.86 (q, J=7.2 Hz, 2H), 2.95 (s, 3H), 3.19 (m, 2H), 4.12 (t, J=8.0 Hz, 2H), 4.79 (br, 1H).

Example 68

Diethyl 2-(N-t-butoxycarbonylaminopropyl)-malonate

20 Sodium hydride (60% in mineral oil, 01.84 g, 46 mmol) suspended in THF (20 mL) at 0°C under argon was carefully treated with diethyl malonate (7 mL, 1.0 equiv) and the mixture was allowed to warm to rt. When it became homogeneous the mesylate **65** (5.84 g, 0.5 equiv) in THF (15 mL) was added all at once. The mixture was heated at reflux overnight, diluted with
25 1:1 hexane/ethyl acetate and washed with water and brine, dried over sodium sulfate and evaporated. The residue was chromatographed on silica, eluting with 25-35% EtOAc/hexane (5.751 g, 79%) ¹H NMR δ 1.18 (t, J=7.9 Hz, 6H), 1.35 (s, 9H), 1.81 (m, 2H), 3.04 (q, J=7.4 Hz, 2H), 3.24 (t, J=6.2 Hz, 1H), 4.10 (q, J=7.9 Hz, 4H), 4.62 (br, 1H). MS (FAB) 318 (MH)⁺,
30 262 (M-C₄H₉), 218 (M-C₅H₉O₂). Anal calcd for C₁₅H₂₇NO₆: C-56.77, H-8.57, N-4.41. Found: C-56.65, H-8.45, N-4.42.

Example 69**N-t-Butoxycarbonyl-5-amino-2-hydroxymethylpentanol**

The diester **66** (4.957 g, 15.62 mmol) in ether (100 mL) under argon at 0°C was treated with lithium borohydride (2M in THF, 24 mL, 3 equiv) and then with methanol (1.9 mL, 3 equiv). The mixture was heated at reflux for 2.5 h and then carefully quenched with methanol (25 mL) followed by acetic acid (2.7 mL, 3 equiv) upon cooling to rt. The mixture was evaporated and flushed with methylene chloride (2x). The residue was dissolved in methanol (100 mL) and the mixture heated at reflux overnight and then evaporated. The resulting oil was dissolved as far as possible in chloroform (100 mL) with sonication and then the solid inorganics were removed by filtration. The filtrate was evaporated and the residue chromatographed on silica, eluting with 9% methanol/methylene chloride, to give the product as a thick oil (2.770 g, 76%) ¹H NMR δ 1.21 and 1.41 (m, each 2H), 1.37 (s, 9H), 1.60 (m, 1H), 2.98 (q, J=7.4 Hz, 2H), 3.68 (m, 4H), 3.96 (br, 1H), 4.97 (brt, 1H). MS (DCI) 234 (MH)⁺, 178 (M-C₄H₉)⁺, 134 (M-C₅H₉O₂)⁺. Anal calcd for C₁₁H₂₃NO₄: C-56.63, H-9.94, N-6.00. Found: C-56.16, H-9.83, N-5.94.

Example 70**5-Amino-2-hydroxymethylpentanol•CF₃CO₂H**

A solution of 5% water in 50% TFA/methylene chloride (4 mL) was added to the diol **67** (0.145 g, 0.621 mmol) and the mixture was stirred for 1 h. The solvents were evaporated and the residue flushed with methylene chloride (2x) (0.153 g, 100%). ¹H NMR δ 1.22 and 1.50 (m, each 2H), 1.69 (m, 1H), 2.82 (br, 2H), 3.51 (d, J=6.9 Hz, 4H), 7.78 (br, 3H). MS (DCI) 134 (MH)⁺.

Example 71**5-Maleimido-2-hydroxymethylpentanol**

The aminodiol **68** in dry acetonitrile under argon at 0°C is treated with maleic anhydride (1 equiv) and DIEA (1 equiv). The mixture is allowed to stir at rt for 4 h. The reaction is re-cooled to 0°C and treated with trimethylsilyl chloride (3 equiv) and DIEA (3 equiv). The mixture is heated at reflux for 4 h and then, after cooling to rt, is evaporated. The residue is partitioned

between ethyl acetate and water. The organic phase is washed with 10% citric acid, water and brine, dried over sodium sulfate and evaporated. The residue is chromatographed on silica gel to give the product.

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Example 72**5-Maleimido-2-hydroxymethylpentanol bis-chloroformate**

A solution of the diol **69** in dry methylene chloride is added dropwise to a vigorously stirred solution of 1M phosgene in toluene (20 equiv) at 0°C. After addition is complete the mixture is stirred at rt for 16 h and then the solvents are removed on the rotary evaporator. The residue is flushed with dry methylene chloride and dried in vacuo. The crude bis-chloroformate is used without further purification.

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Example 73**MPr-BHP-(Phe-Lys(MMT)-PABOH)₂**

To a stirred solution of the crude bis-chloroformate **70** in dry methylene chloride at 0°C is added a solution of Phe-Lys(MMT)-PABOH **10a** (2 equiv) and DIEA (2 equiv) in methylene chloride all at once. The mixture is allowed to stir at rt for 16 h and then the solvent is evaporated. The residue is partitioned between ethyl acetate and water. The organic phase is washed with brine, dried over sodium sulfate and evaporated. The product is purified by column chromatography using silica gel.

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Example 74

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MPr-BHP-(Phe-Lys(MMT)-PABC-PNP)₂

A stirred mixture of **71**, bis-p-nitrophenylcarbonate (6 equiv) and freshly-activated 4Å powdered sieves in dry methylene chloride under argon at rt is treated with DIEA (6 equiv). After continued stirring for 16 h the mixture is filtered and the filtrate evaporated. The residue is dried in vacuo for several hours and then dissolved in a minimum volume of methylene chloride. To this is added two-times the volume of ether. The resulting solid is collected by filtration, washed with 2:1 ether/methylene chloride, and dried in vacuo.

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Example 75**MPr-BHP-(Phe-Lys(MMT)-PABC-DOX)₂**

A stirred solution of the bis-p-nitrophenylcarbonate **72** and DOX hydrochloride (1.1 equiv) in DMF is treated with DIEA (1.1 equiv). After stirring for 2 d the mixture is diluted with ethyl acetate. The solution is washed with water (4x) and brine, dried over sodium sulfate and evaporated. The residue is chromatographed on silica gel to give the product.

Example 76**MPr-BHP-(Phe-Lys-PABC-DOX)₂•2Cl₂CHCO₂H**

A stirred solution of **73** in methylene chloride is treated with anisole (100 equiv) and then dichloroacetic acid (10 equiv). After 1.5 h the mixture is diluted with ethyl acetate. The resulting suspension is stirred at rt for 1 h and then the orange solid product is collected by filtration, washed with ethyl acetate and dried in vacuo.

Example 77**N-Adpoc-3-hydroxyaminopropane**

A stirred solution of 3-hydroxyaminopropane and sodium bicarbonate (1 equiv) in water at rt is treated with 1-(1'-adamantyl)-1-methyl-ethoxycarbonyl chloride (1 equiv). After 6 h the resulting solid product is collected by filtration, washed with water, and lyophilized in vacuo. The crude product is carried on without further purification.

Example 78**N-Adpoc-3-methanesulfonyloxyaminopropane**

To a stirred solution of the alcohol **75** in methylene chloride at 0°C is added methanesulfonyl chloride (1.05 equiv) and then triethylamine (1.05 equiv). The mixture is allowed to warm to rt and stirred overnight at rt. The solvents is evaporated and the residue partitioned between ethyl acetate and pH 5 buffer (bipthalate). The organic phase is washed with water and brine, dried over sodium sulfate and evaporated to give the product which is carried on without further purification.

Example 79**t-Butyl N-adpoc-5-amino-2-t-butoxycarbonyl-pentanoate**

A stirred suspension of sodium hydride in dry tetrahydrofuran at 0°C under argon is carefully treated with di-t-butylmalonate (1 equiv). The mixture is gradually allowed to warm to rt. When the mixture is homogeneous it is re-cooled to 0°C and then treated all at once with a solution of the mesylate **76** (0.8 equiv) in tetrahydrofuran. The mixture is allowed to warm to rt and then is heated at reflux for 16 h. Upon cooling to rt the mixture is diluted with ether and the solution is washed with sat. ammonium chloride, water and brine, dried over sodium sulfate and evaporated. The residue is chromatographed on silica gel to give the product.

Example 80**t-Butyl 5-amino-2-t-butoxycarbonyl-pentanoate**

The Adpoc substrate **77** is dissolved in 5% trifluoroacetic acid/methylene chloride containing anisole (100 equiv). After 30 min the solvents are removed under high vacuum at rt and the residue dried in vacuo for several hours. The crude product is carried on immediately without further purification.

Example 81**t-Butyl 5-maleimido-2-t-butoxycarbonyl-pentanoate 79**

The aminodiester **78** in dry acetonitrile under argon at 0°C is treated with maleic anhydride (1 equiv) and the stirred mixture is allowed to stir at rt for 4 h. The mixture is re-cooled to 0°C and treated with trimethylsilyl chloride (3 equiv) and DIEA (3 equiv). The mixture is heated at reflux for 4 h and then, after cooling to rt, is evaporated. The residue is partitioned between ethyl acetate and water. The organic phase is washed with 10% citric acid, water and brine, dried over sodium sulfate and evaporated. The residue is chromatographed on silica to give the product.

Example 82**2-(3'-maleimidopropyl)-malonic acid**

The di-t-butyl ester **79** is dissolved in 20% trifluoroacetic acid and the mixture is stirred at rt for 5 h. The solvents are evaporated and the residue
5 flushed with methylene chloride several times. The residue is triturated with ether and the resulting solid product is collected by filtration, washed with ether, and dried in vacuo.

Example 83**MPr-Mal-(Phe-Lys(MMT)-PABOH)₂**

A stirred solution of the diacid **80** in dimethoxyethane at 0°C is treated with n-hydroxysuccinimide (2.2 equiv) and 0.5 M DCC in methylene chloride (2.2 equiv). The mixture is allowed to warm to rt. After 4 h a solution of Phe-Lys(MMT)-PABOH **10a** (2 equiv) in dimethoxyethane is added and the
15 mixture is stirred at rt for 16 h. The mixture is then filtered and the filtrate evaporated. The residue is partitioned between ethyl acetate and water. The organic phase is washed with water and brine, dried over sodium sulfate and evaporated. The product is purified by chromatography on silica gel.

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Example 84**MPr-Mal-(Phe-Lys(MMT)-PABC-PNP)₂**

A stirred mixture of **81**, bis-p-nitrophenylcarbonate (6 equiv) and freshly activated 4Å powdered sieves in dry methylene chloride under argon at rt is
25 treated with DIEA (6 equiv). After continued stirring for 16 h the mixture is filtered and the filtrate evaporated. The residue is dried in vacuo for several hours and then dissolved in a minimum volume of methylene chloride. To this is added two-times the volume of ether. The resulting solid is collected by filtration, washed with 2:1 ether/methylene chloride, and dried in vacuo.

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Example 85**MPr-Mal-(Phe-Lys(MMT)-PABC-DOX)₂**

A stirred solution of the bis-p-nitrophenylcarbonate **82** and DOX hydrochloride (1.1 equiv) in DMF is treated with DIEA (1.1 equiv). After

stirring for 2 d the mixture is diluted with ethyl acetate. The solution is washed with water (4x) and brine, dried over sodium sulfate and evaporated. The residue is chromatographed on silica gel to give the product.

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Example 86**MPr-Mal-(Phe-Lys-PABC-DOX)₂•2Cl₂CHCO₂H**

A stirred solution of **83** in methylene chloride is treated with anisole (100 equiv) and then dichloroacetic acid (10 equiv). After 1.5 h the mixture is diluted with ethyl acetate. The resulting suspension is stirred at rt for 1 h and then the orange solid product is collected by filtration, washed with ethyl acetate and dried in vacuo.

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Example 87**N-Carbobenzyloxy-N', N'-bis-(carboxymethyl)-**

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ethylenediamine

The di-t-butyl ester **22** is dissolved in 20% trifluoroacetic acid and the mixture is stirred at rt for 5 h. The solvents are evaporated and the residue flushed with methylene chloride several times. The residue is triturated with ether and the resulting solid product is collected by filtration, washed with ether, and dried in vacuo.

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Example 88**N-Carbobenzyloxy-N', N'-bis-(2-hydroxyethyl)-****ethylenediamine 86**

A solution of the bis-carboxylic acid **85** in diglyme at 0°C is treated with N-hydroxysuccinimide (2.2 equiv) and DCC (2.2 equiv). The mixture is allowed to warm to rt and then stirred for 16 h. The resulting solid is removed by filtration. The filtrate is cooled to 0°C under an argon atmosphere and then treated with 0.5 M sodium borohydride in diglyme. The mixture is allowed to warm to rt for 16 h and is then poured into ice water. The resulting mixture is extracted with ethyl acetate. The organic phase is washed with water and brine, dried over sodium sulfate and evaporated to give the product which is carried on without further purification.

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Example 89**Bis-(2-hydroxyethyl)ethylenediamine hydrochloride**

A solution of diol **86** and conc. HCl (1 equiv) in methanol is degassed by
5 bubbling nitrogen through it for 30 min. To this is added 10% palladized
charcoal (5 wt. percent). The resulting mixture is shaken on a Parr
apparatus under 50 psi of hydrogen gas for 16 h. The mixture is then filtered
and the filtrate evaporated. The residue is flushed several times with
chloroform to drive off traces of methanol. The crude product is carried on
10 without further purification.

Example 90**Bis-(2-hydroxyethyl)-2'-maleimidoethylamine**

The aminodiol **87** in dry acetonitrile under argon at 0°C is treated with
15 maleic anhydride (1 equiv) and DIEA (1 equiv). The mixture is allowed to stir
at rt for 4 h. The mixture is re-cooled to 0°C and treated with trimethylsilyl
chloride (3 equiv) and DIEA (3 equiv). The reaction is heated at reflux for 4 h
and then, after cooling to rt, is evaporated. The residue is partitioned
between ethyl acetate and water. The organic phase is washed with 10%
20 citric acid, water and brine, dried over sodium sulfate and evaporated. The
residue is chromatographed on silica gel to give the product.

Example 91**Bis-(2-chlorocarbonyloxyethyl)-2'-maleimidoethylamine**

25 A solution of the diol **88** in dry methylene chloride is added dropwise to a
vigorously stirred solution of 1M phosgene in toluene (20 equiv) at 0°C.
After addition is complete the mixture is stirred at rt for 16 h and then the
solvents are removed on the rotary evaporator. The residue is flushed with
dry methylene chloride and dried in vacuo. The crude bis-chloroformate is
30 used without further purification.

Example 92**MEt-IBHE-(Phe-Lys(MMT)-PABOH)₂**

To a stirred solution of the crude bis-chloroformate **89** in dry methylene chloride at 0°C is added a solution of Phe-Lys(MMT)-PABOH **10a** (2 equiv) and DIEA (2 equiv) in methylene chloride all at once. The mixture is allowed to stir at rt for 16 h and then the solvent is evaporated. The residue is partitioned between ethyl acetate and water. The organic phase is washed with brine, dried over sodium sulfate and evaporated. The product is purified by column chromatography using silica gel.

Example 93**MEt-IBHE-(Phe-Lys(MMT)-PABC-PNP)₂**

A stirred mixture of **90**, bis-p-nitrophenylcarbonate (6 equiv) and freshly activated 4Å powdered sieves in dry methylene chloride under argon at rt is treated with DIEA (6 equiv). After continued stirring for 16 h the mixture is filtered and the filtrate evaporated. The residue is dried in vacuo for several hours and then dissolved in a minimum volume of methylene chloride. To this is added two-times the volume of ether. The resulting solid is collected by filtration, washed with 2:1 ether/methylene chloride, and dried in vacuo.

Example 94**MEt-IBHE-(Phe-Lys(MMT)-PABC-DOX)₂**

A stirred solution of the bis-p-nitrophenylcarbonate **91** and DOX hydrochloride (1.1 equiv) in DMF is treated with DIEA (1.1 equiv). After stirring for 2 d the mixture is diluted with ethyl acetate. The solution is washed with water (4x) and brine, dried over sodium sulfate and evaporated. The residue is chromatographed on silica gel to give the product.

Example 95**MEt-IBHE-(Phe-Lys-PABC-DOX)₂•2Cl₂CHCO₂H**

A stirred solution of **92** in methylene chloride is treated with anisole (100 equiv) and then dichloroacetic acid (10 equiv). After 1.5 h the mixture is diluted with ethyl acetate. The resulting suspension is stirred at rt for 1 h and

then the orange solid product is collected by filtration, washed with ethyl acetate and dried in vacuo.

Example 96

5 114 (257 mg, 0.25 mmole) and Doxorubicin HCl (377 mg, 0.65 mmole) are stirred in 25 ml dry methanol for 24 hour. The reaction is concentrated by rotary evaporation to 4 ml, then purified in two equal portions on Sephadex LH-20 (1" x 18") with methanol. Fractions containing pure product are pooled, rotary evaporated, and dried under high vacuum to yield 115 (222
10 mg, 50%).

Example 97

Conjugation of BR96 or IgG and Branched Peptide Linker

Thiolation:

15 BR96 and IgG were thiolated (reduced) by a variation of previously reported method¹. In a typical example, 1.27 g BR96 (75 ml at 105.7 uM, 7.93 umole) was de-oxygenated by several cycles of alternating vacuum and Ar atmosphere. This was then treated with 7.4 mM DTT (8.6 ml, 63.6 umole in Ar-bubbled PBS, pH 7.0) and stirred at 37°C under Ar for 3 hr. Removal of
20 low molecular weight compounds was accomplished by ultrafiltration against PBS, pH 6.0 in an Amicon stirred cell at 4°C. A 400 ml Amicon cell was fitted with an Amicon YM30 filter (molecular weight cut-off 30,000), and charged to 40 psi with Ar. Cell eluant was monitored for thiol content with Ellman's reagent² until a baseline reading at 412 nm was obtained. Concentration of
25 protein and thiol groups were determined according to the previously reported method¹. In this example, 1.10 g reduced BR96 (85 ml at 80.8 uM MAb, 619.1 uM thiol) was obtained, for a yield of 87% and a thiol titer of 7.7 mole thiol groups/mole BR96.

30

Example 98

Conjugation:

The following procedure,¹ for the conjugation of BR96 and M-Et-IDA-(Phe-Lys-PABC-DOX= β -Ala-mTEG)₂, is typical of that used for all branched

peptide linkers modified with the β -Ala-mTEG hydrazone or β -Ala
hydrazone. To reduced BR96 (83 ml, 6.70 μ mole MAb, 51.6 μ mole thiol) was
added dropwise at 0°C under Ar a solution of M-Et-IDA-(Phe-Lys-PABC-
DOX= β -Ala-mTEG)₂ (239 mg, 77.3 μ mole) in 4.0 ml DMSO. After stirring for
5 30 min., the reaction was filtered through a 0.22 μ sterile filter. Conjugate was
purified at 4°C by percolation (approximately 2 ml/min.) through a 1" x 24"
Bio-Beads column (initially prepared by swelling and packing in methanol,
then equilibrated in H₂O, and finally PBS, pH 7.4). The purified conjugate
was filtered again through a 0.22 μ sterile filter to yield 95 ml of BR96-S-Et-
10 IDA-(Phe-Lys-PABC-DOX= β -Ala-mTEG)₂ (BR96, 76.1 μ M; DOX, 1.11 mM;
MR, 14.6 mole DOX/mole BR96; yield, 100%). Conjugate was frozen in liquid
N₂ and stored at -80°C.

For more guidance, see Willner, D., Trail, P. A., Hofstead, S. J., King,
H. D., Lasch, Braslawsky, G. R., Greenfield, R. S., Kaneko, T., Firestone, R. A.
15 (1993) (6-Maleimidocaproyl)hydrazone of Doxorubicin: A new derivative for
the preparation of immunoconjugates of Doxorubicin. Bioconjugate Chem.,
4, 521 and Riddles, P. W., Blakeley, R. L., Zerner, B., (1979) Ellman's
reagent: 5,5'-Dithiobis(2-nitrobenzoic acid)-A reexamination. Anal.
Biochem., 94, 75.

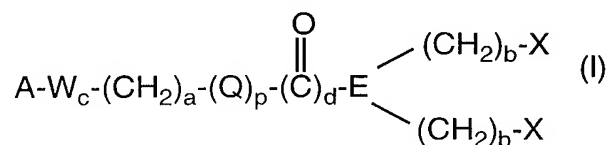
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What is Claimed is:

1. A branched peptide linker for linking a thiol group derived from a targeting ligand to two or more drug moieties which comprises a compound having a terminus containing a thiol acceptor for binding to a thiol group
5 derived from a targeting ligand, at least one point of branching which is a polyvalent atom allowing for a level of branching of 2^n , wherein n is a positive integer, at least two amino acid moieties providing an enzymatic cleavage site, and at least two other termini containing groups capable of forming covalent bonds with chemically reactive functional groups derived
10 from a drug moiety.
2. The branched peptide linker of Claim 1 wherein n is 1, 2, 3, or 4.
3. The branched peptide linker of Claim 1 wherein n is 1, 2, or 3.
15
4. The branched peptide linker of Claim 1 wherein n is 1 or 2.
5. The branched peptide linker of Claim 1 linked to an antibody or fragment thereof.
20
6. The branched peptide linker of Claim 1 wherein said polyvalent atom is nitrogen or carbon.
7. The branched peptide linker of Claim 1 wherein said polyvalent
25 is nitrogen.
8. The branched peptide linker of Claim 1 wherein said polyvalent is carbon.
9. The branched peptide linker of Claim 1 linked to drug moieties which are anthracyclines.
30

10. The branched peptide linker of Claim 1 wherein n is 1 or 2, said targeting ligand is an antibody or fragment thereof, said polyvalent atom is nitrogen and said drug moieties are anthracyclines.

5 11. A compound of the formula



wherein

A is a thiol acceptor;

W is a bridging moiety;

10 c is an integer of 0 to 1;

a is an integer of 2 to 12;

Q is O, NH, or N-lower alkyl;

p is an integer of 0 or 1;

d is an integer of 0 or 1;

15 E is a polyvalent atom;

each b is an integer of 1 to 10;

each X is of the formula

-CO-Y-Z_m-G_n

wherein

20 Y is two amino acid residues in the L form;

Z is one or two amino acid residues;

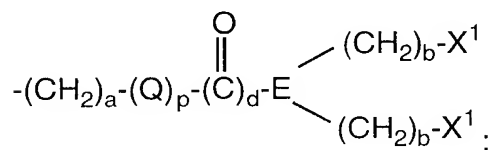
m is an integer of 0 or 1;

G is a self-immolative spacer; and

n is a integer of 0 or 1; provided that when G is 0 then -Y-Z_m

25 is ala-leu-ala-leu or gly-phe-leu-gly;

or each X is of the formula

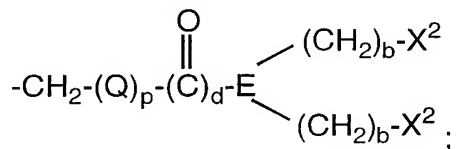


wherein each X¹ is of the formula

-CO-Y-Z_m-G_n;

and wherein Y, Z, Q, E, G, m, d, p, a, b and n are as defined above;

or each X¹ is of the formula



5

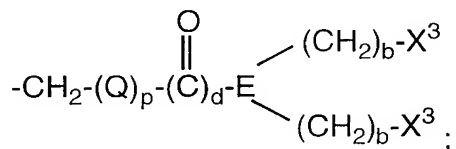
wherein each X² is of the formula

-CO-Y-Z_m-G_n;

and wherein Y, Z, G, Q, E, m, d, p, a, b and n are as defined above;

10

or each X² is of the formula



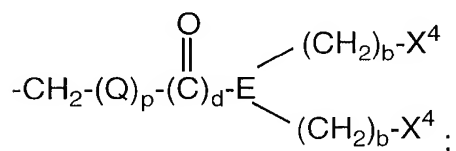
wherein each X³ is of the formula

-CO-Y-Z_m-G_n;

and wherein Y, Z, G, Q, E, m, d, p, a, b and n are as defined above;

15

or each X³ is of the formula



wherein each X⁴ is of the formula

-CO-Y-Z_m-G_n;

20

and wherein Y, Z, G, Q, E, m, d, p, a, b and n are as defined above.

12. The compound of Claim 11 wherein each X is of the formula

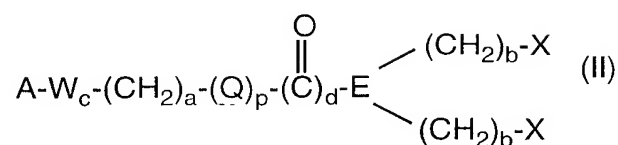
-CO-Y-Z_m-G_n

wherein Y, m, Z, and n are as defined above.

25

13. The compound of Claim 11 wherein each X^1 is of the formula $\text{—CO—Y—Z}_m\text{—G}_n$ wherein Y, m, Z, and n are as defined above.
- 5 14. The compound of Claim 11 wherein each X^2 is of the formula $\text{—CO—Y—Z}_m\text{—G}_n$ wherein Y, m, Z, and n are as defined above.
- 10 15. The compound of Claim 11 wherein each X^3 is of the formula $\text{—CO—Y—Z}_m\text{—G}_n$ wherein Y, m, Z, and n are as defined above.
- 15 16. The compound of Claim 11 wherein —Y—Z_m is a dipeptide, tripeptide or tetrapeptide containing amino acids selected from the group consisting of alanine, valine, leucine, isoleucine, methionine, phenylalanine, tryptophan, proline, lysine, lysine protected with acetyl or formyl, arginine, arginine protected with tosyl or nitro groups, histidine, ornithine, ornithine protected with acetyl or formyl, and citrulline.
- 20 17. The compound of Claim 11 wherein the —Y—Z_m group is phenylalanine-lysine.
- 25 18. The compound of Claim 11 wherein the —Y—Z_m group is valine-citrulline.
19. The compound of Claim 11 wherein the —Y—Z_m group is valine-lysine.
20. The compound of Claim 11 wherein E is nitrogen.
- 30 21. The compound of Claim 11 wherein E is carbon.

22. A compound of the formula



wherein

A is a thiol acceptor;

5 W is a bridging moiety;

c is an integer of 0 to 1;

a is an integer of 2 to 12;

Q is O, NH, or N-lower alkyl;

p is an integer of 0 or 1;

10 d is an integer of 0 or 1;

E is a polyvalent atom;

each b is an integer of 1 to 10;

each X is of the formula

-CO-Y-Z_m-G_nD

15 wherein

Y is two amino acid residues in the L form;

Z is one or two amino acid residues;

m is an integer of 0 or 1;

G is a self-immolative spacer;

20 n is a integer of 0 or 1; provided that when G is 0 then -Y-Z_m-

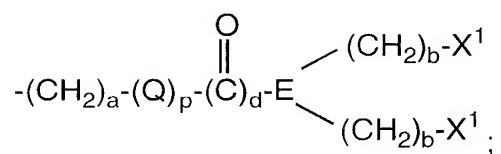
is ala-leu-ala-leu or gly-phe-leu-gly; and

D is a Drug moiety having a backbone and at least one chemically reactive functional group pendant thereto

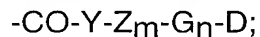
chemically

25 reacted to the self-immolative spacer or terminal amino acid residue to form a covalent bond, said functional group selected from the group consisting of a primary or secondary amine, hydroxyl, sulfhydryl, carboxyl, aldehyde or ketone;

or each X is of the formula

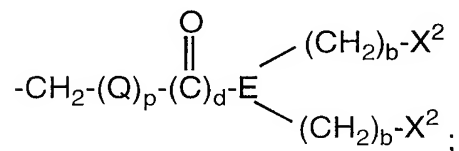


wherein each X^1 is of the formula

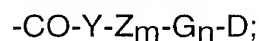


wherein Y, Z, G, D, Q, E, m, d, p and n are as defined above;

5 or each X^1 is of the formula



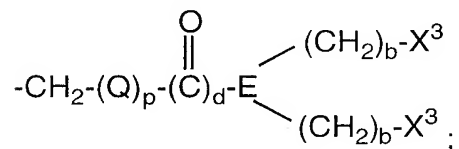
wherein each X^2 is of the formula



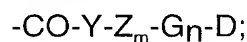
and wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined
above;

10

or each X^2 is of the formula

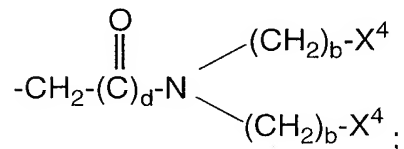


wherein each X^3 is of the formula

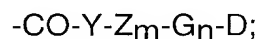


15 and wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined
above;

or each X^3 is of the formula



and wherein each X^4 is of the formula



20

and wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined
above.

23. The compound of Claim 22 wherein each X is of the formula
-CO-Y-Z_m-G_n-D
wherein Y, m, Z, n, and D are as defined above.

5

24. The compound of Claim 22 wherein each X¹ is of the formula
-CO-Y-Z_m-G_n-D
wherein Y, m, Z, n, and D are as defined above.

10 25. The compound of Claim 22 wherein each X² is of the formula
CO-Y-Z_m-G_n-D
wherein Y, m, Z, n, and D are as defined above.

15 26. The compound of Claim 22 wherein each X³ is of the formula
-CO-Y-Z_m-G_n-D
wherein Y, m, Z, n, and D are as defined above.

27. The compound of Claim 22 wherein -Y-Z_m is a dipeptide,
tripeptide or tetrapeptide containing amino acids selected from the group
20 consisting of alanine, valine, leucine, isoleucine, methionine, phenylalanine,
tryptophan, proline, lysine, lysine protected with acetyl or formyl, arginine,
arginine protected with tosyl or nitro groups, histidine, ornithine, ornithine
protected with acetyl or formyl, and citrulline.

25 28. The compound of Claim 17 wherein the -Y-Z_m group is
phenylalanine-lysine.

29. The compound of Claim 17 wherein the -Y_m-Z_m group is
valine-citrulline.

30

30. The compound of Claim 17 wherein the -Y-Z_m group is valine-
lysine.

31. The compound of Claim 22 wherein E is nitrogen.

32. The compound of Claim 22 wherein E is carbon.

5

33. The compound of Claim 22 wherein D is an amino containing drug moiety selected from the group consisting of mitomycin-C, mitomycin-A, daunorubicin, doxorubicin, N-(5,5-diacetoxypentyl)doxorubicin, aminopterin, actinomycin, bleomycin, 9-amino camptothecin, N8-acetyl spermidine, 1-(2
10 chloroethyl)-1,2-dimethanesulfonyl hydrazide, tallysomycin, and derivatives thereof.

34. The compound of Claim 22 wherein D is doxorubicin.

15

35. The compound of Claim 22 wherein D is mitomycin-C.

36. The compound of Claim 22 wherein D is mitomycin-A.

37. The compound of Claim 22 wherein D is tallysomycin.

20

38. The compound of Claim 22 wherein D is N-(5,5-diacetoxypentyl)doxorubicin.

39. The compound of Claim 22 wherein D is a hydroxyl containing
25 drug moiety selected from the group consisting of etoposide, camptothecin, taxol, esperamicin, 1,8-dihydroxy-bicyclo[7.3.1]trideca-4-ene-2,6-diyne-13-one, anguidine, doxorubicin, morpholine-doxorubicin, N-(5,5-diacetoxypentyl)doxorubicin, vincristine, vinblastine and derivatives thereof.

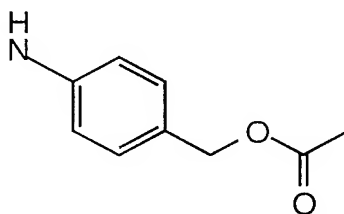
30 40. The compound of Claim 22 wherein D is a sulfhydryl containing drug moiety selected from the group consisting of esperamicin and 6-mercaptopurine, and derivatives thereof.

41. The compound of Claim 22 wherein D is a carboxyl containing — drug moiety selected from the group consisting of methotrexate, camptothecin (ring-opened form of the lacton), butyric acid, retinoic acid, and derivatives thereof.

5

42. The compound of Claim 22 wherein D is an anthracycline.

43. The compound of Claim 22 wherein G is

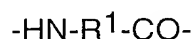


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44. The compound of Claim 22 wherein G is p-aminobenzyl-carbonyl.

15

45. The compound of Claim 22 wherein G has the formula



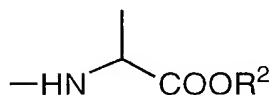
wherein

20

R^1 is C_1 - C_5 alkyl,

T is O, N or S.

46. The compound of Claim 22 wherein G has the formula

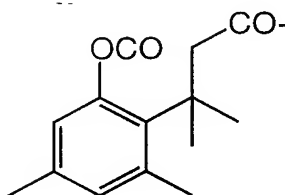


25

wherein

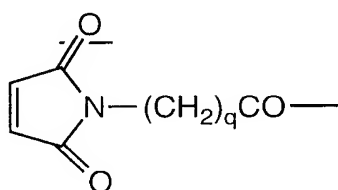
R^2 is H or C_1 - C_5 alkyl.

47. The compound of Claim 22 wherein G is



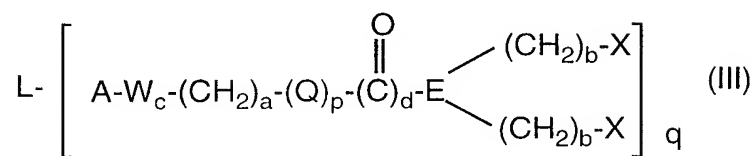
5

48. The compound of Claim 22 wherein A is



10

49. A compound of the formula



wherein

L is a ligand;

q is an integer of 1 to 10;

A is a thiol acceptor;

W is a bridging moiety;

c is an integer from 0 to 1;

a is an integer of 2 to 12;

Q is O, NH, or N-lower alkyl;

p is an integer of 0 or 1;

d is an integer of 1 or 2;

E is a polyvalent atom;

each b is an integer of 1 to 10;

each X is of the formula

15

20

-CO-Y-Z_m-G_n-D

wherein

Y is two amino acid residues in the L form;

Z is one or two amino acid residues;

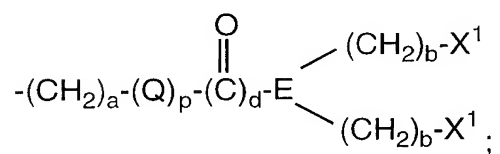
5 m is an integer of 0 or 1;

G is a self-immolative spacer; and

n is a integer of 0 or 1; provided that when G is 0 then -Y-Z_m- is ala-leu-ala-leu or gly-phe-leu-gly;

10 D is a drug moiety having a backbone and at least one chemically reactive functional group pendant thereto reacted to the self-immolative spacer to form a covalent bond, said functional group selected from the group consisting of a primary or secondary amine, hydroxyl, carboxyl, sulfhydryl, aldehyde, or ketone;

15 or X is of the formula

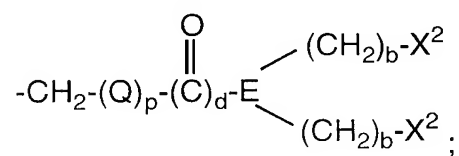


wherein each X¹ is of the formula

-CO-Y-Z_m-G_n-D;

20 and wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined above;

or each X¹ is of the formula

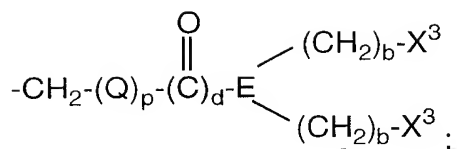


wherein each X² is of the formula

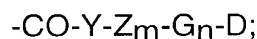
-CO-Y-Z_m-G_n-D;

25 and wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined above;

or each X² is of the formula



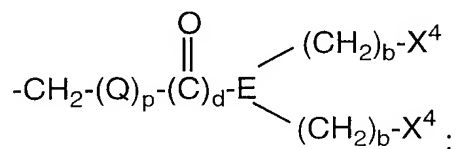
wherein each X^3 is of the formula



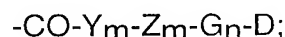
wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined
above;

5

or each X^3 is of the formula



wherein end X^4 is of the formula



10

wherein Y, Z, G, D, Q, E, m, d, p, a, b and n are as defined
above.

50. The compound of Claim 49 wherein L is an immunoglobulin, or
a fragment thereof.

15

51. The compound of claim 49 wherein L is an immunoglobulin
selected from the group consisting of BR96, BR64, L6, a relaxed BR96, a
relaxed BR64, a chimeric L6, a relaxed chimeric BR96, a relaxed chimeric
BR64, a relaxed chimeric L6; and, fragments thereof.

20

52. The compound of claim 49 wherein L is a ligand selected from
the group consisting of bombesin, EDG, transferrin, gastrin, gastrin-releasing
peptide, platelet-derived growth factor, IL-2, IL-6, TFG- α , TFG- β , VGF, insulin,
insulin-like growth factors I and II, carbohydrates, lectins, and apoproteins
from low density lipoproteins.

25

53. A pharmaceutical composition comprising a compound of
Claim 49 and a pharmaceutically acceptable carrier.

54. A method of controlling the growth of undesirable cells which comprises administering a compound of Claim 49 to a host in need of such treatment.

5

55. The method of claim 54 wherein the undesirable cells are cancer cells.

56. A method of treatment of a warm blooded animal in need thereof which comprises administering to said animal an effective tumor inhibiting amount of a compound of Claim 49.

10

57. A method of targeting a pharmaceutical agent to a desired site in a host or sample comprising administering to the host or contacting the sample with a compound of Claim 49.

15

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/19851

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : Please See Extra Sheet.

US CL : 530/350, 333; 424/130.1, 141.1, 136.1, 144.1

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 530/350, 333; 424/130.1, 141.1, 136.1, 144.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, DIALOG, SEARCH TERMS: PEPTIDE LINKER, CHIMERIC ANTIBODIES, BRANCHED PEPTIDES, THIOETHER CONJUGATES, IMMUNO CONJUGATES

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 91/18016 A1 (DEGHENHGI) 28 November 1991, entire document.	1-57
Y	EP 0 328 147 A2 (BRISTOL-MYERS COMPANY) 16 August 1989, pages 1-29.	1-57
Y	EP 0 457 250 A2 (BRISTOL-MYERS SQUIBB COMPANY) 21 November 1991, pages 24.	1-57
Y, P	US 5,622,929 A (WILLNER ET AL) 22 April 1997, entire document.	1-57
Y	US 4,975,278 A (SETER ET AL) 04 December 1990, entire document.	1-57
Y	US 5,122,368 A (GREENFIELD ET AL) 16 June 1992, entire document.	1-57

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
A document defining the general state of the art which is not considered to be of particular relevance	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
E earlier document published on or after the international filing date	*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/19851

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y, P	US 5,606,017 A (WILLNER ET AL) 25 February 1997, entire document.	1-57
Y	US 5,563,250 A (HYALARIDES ET AL) 08 October 1996, entire document.	1-57
Y	US 5,349,066 A (KANEKO ET AL) 20 September 1994, entire document.	1-57
Y	US 5,137,877 A (KANEKO ET AL) 11 August 1992, entire document.	1-57
Y	US 5,017,693 A (HYALRIDES ET AL) 21 May 1991, entire document.	1-57
Y	US 5,084,560 A (HELLSTROM ET AL) 28 January 1992, entire document.	1-57
Y	US 5,116,944 A (SIVAM ET AL) 26 May 1992, entire document.	1-57
Y, P	US 5,635,603 A (HANSEN ET AL) 03 June 1997, entire document.	1-57
Y	US 5,252,713 A (MORGAN, JR. ET AL) 12 October 1993, entire document.	1-57
Y	US 5,489,516 A (BROUDY ET AL) 06 February 1996, entire document.	1-57

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A. CLASSIFICATION OF SUBJECT MATTER:

IPC (6):

A61K 39/395, 39/40, 39/42, 38/00; C07K 1/00, 14/00, 17/00, 5/00, 7/00, 16/00